



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<b>(21) International Application Number:</b> PCT/EP92/00002 <b>(22) International Filing Date:</b> 2 January 1992 (02.01.92)  <b>(30) Priority data:</b> 91100123.8 3 January 1991 (03.01.91) EP <b>(34) Countries for which the regional or international application was filed:</b> AT et al.  <b>(71) Applicant (for all designated States except US):</b> GRUPPO LEPETIT S.P.A. [IT/IT]; Via Murat, 23, I-20159 Milano (IT).  <b>(72) Inventors; and</b> <b>(75) Inventors/Applicants (for US only):</b> TAVECCHIA, Paolo [IT/IT]; Via Buon Gesù, 1, I-20017 Rho (IT). LOCIURO, Sergio [IT/IT]; Via Troilo, 4, I-20136 Milano (IT). CIABATTI, Romeo [IT/IT]; Via Brodolini, 15/A, I-20026 Novate Milanese (IT). SELVA, Enrico [IT/IT]; Via Di Vittorio, 15, I-27027 Gropello Cairoli (IT).	<b>(74) Agent:</b> PIMPINELLI, Natale; Gruppo Lepetit S.p.A., Patent & Trademark Department, Via Roberto Lepetit, 34, I-21040 Gerenzano (IT).  <b>(81) Designated States:</b> AT (European patent), AU, BE (European patent), CA, CH (European patent), DE (European patent), DK (European patent), ES (European patent), FI, FR (European patent), GB (European patent), GR (European patent), HU, IT (European patent), JP, KR, LU (European patent), MC (European patent), NL (European patent), NO, SE (European patent), US.  <b>Published</b> <i>With international search report.</i>	
<b>(54) Title:</b> AMIDES OF ANTIBIOTIC GE 2270 FACTORS		
<b>(57) Abstract</b>  The present invention is directed to novel amide derivatives of antibiotic GE 2270 compounds and a process for preparing them. Said amide derivatives are antimicrobial agents active against gram positive bacteria as well as gram negative bacteria.		

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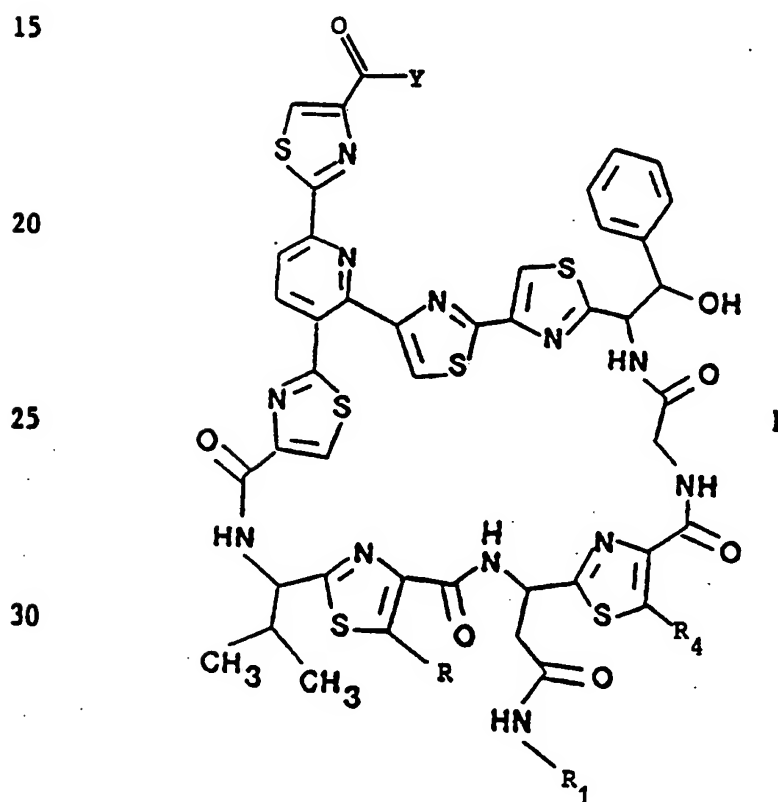
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## AMIDES OF ANTIBIOTIC GE 2270 FACTORS

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10           The present invention is directed to novel amide derivatives of antibiotic GE 2270 having the following formula I



wherein

R represents:  
hydrogen,  
hydroxymethyl, or  
methoxymethyl;

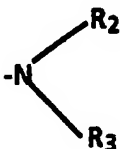
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R<sub>1</sub> represents:  
hydrogen, or  
methyl;

10

Y represents:  
a group of formula

15



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wherein:

R<sub>2</sub> represents:  
hydrogen,  
(C<sub>1</sub>-C<sub>4</sub>)alkyl,  
amino(C<sub>2</sub>-C<sub>4</sub>)alkyl,  
(C<sub>1</sub>-C<sub>4</sub>)alkylamino-(C<sub>1</sub>-C<sub>4</sub>)alkyl, or  
di-(C<sub>1</sub>-C<sub>4</sub>)alkylamino-(C<sub>1</sub>-C<sub>4</sub>)alkyl;

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R<sub>3</sub> represents:  
hydrogen,  
a linear or branched (C<sub>1</sub>-C<sub>14</sub>)alkyl group bearing  
from 1 to 3 substituents selected from: carboxy,



5 sulfo, phosphono, amino which may be optionally  
protected with a lower alkoxy carbonyl or a  
benzyloxy carbonyl group, (C<sub>1</sub>-C<sub>4</sub>)alkylamino  
wherein the alkyl moiety may be optionally  
substituted with a carboxy group,  
10 di-(C<sub>1</sub>-C<sub>4</sub>)alkylamino, hydroxy, halo,  
(C<sub>1</sub>-C<sub>4</sub>)alkoxy wherein the alkyl moiety may be  
optionally substituted with a carboxy group,  
(C<sub>1</sub>-C<sub>4</sub>)alkoxy carbonyl, mercapto,  
15 (C<sub>1</sub>-C<sub>4</sub>)alkylthio wherein the alkyl moiety may be  
optionally substituted with a carboxy group,  
phenyl which may be optionally substituted with  
1 to 3 substituents selected from carboxy,  
20 sulfo, hydroxy, halo and mercapto, carbamyl,  
(C<sub>1</sub>-C<sub>6</sub>)alkylcarbamyl wherein the alkyl moiety  
may be optionally substituted with 1 or 2  
substituents selected from carboxy, amino,  
(C<sub>1</sub>-C<sub>4</sub>)alkylamino and di-(C<sub>1</sub>-C<sub>4</sub>)alkylamino,  
25 di-(C<sub>1</sub>-C<sub>4</sub>)alkylcarbamyl wherein the alkyl  
moieties together with the adjacent nitrogen  
atom may also represent a saturated 5-7 membered  
heterocyclic ring which may optionally be  
substituted with a carboxy or a carbamyl group  
25 on one of the ring carbons and may optionally  
contain a further heterogroup selected from O, S  
and N, benzoylamino wherein the phenyl group may  
be substituted from 1 to 3 hydroxy group, a  
30 nitrogen containing 5-6 membered heterocyclic  
ring which may be unsaturated, partially  
saturated or wholly saturated and may contain 1  
to 3 further heteroatoms selected from N, S and  
O wherein one of the carbons of the ring may  
optionally bear a group carboxy, sulfo,  
carboxy(C<sub>1</sub>-C<sub>4</sub>)alkyl and sulfo(C<sub>1</sub>-C<sub>4</sub>)alkyl and

the ring nitrogen atom may optionally be substituted by (C<sub>1</sub>-C<sub>4</sub>)alkyl, carboxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, sulfo(C<sub>1</sub>-C<sub>4</sub>)alkyl, and benzyl;  
5 (C<sub>3</sub>-C<sub>6</sub>)alkenyl, optionally substituted by carboxy or sulfo;  
1-deoxy-1-glucityl;  
2-deoxy-2-glucosyl;  
10 a fully saturated 5 to 7 membered nitrogen containing heterocyclic ring wherein the nitrogen atom may be optionally substituted by (C<sub>1</sub>-C<sub>4</sub>)alkyl or benzyl and one or two carbons of the ring skeleton may bear a substituent  
15 selected from (C<sub>1</sub>-C<sub>4</sub>)alkyl, carboxy and sulfo;

or R<sub>2</sub> and R<sub>3</sub>

20 taken together with the adjacent nitrogen atom represent a fully saturated 5-7 membered heterocyclic ring which may optionally contain a further heteroatom selected from O, S and N, and may optionally bear one or two substituents on  
25 the ring carbons selected from (C<sub>1</sub>-C<sub>4</sub>)alkyl, benzyl, carboxy, sulfo, carboxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, and sulfo(C<sub>1</sub>-C<sub>4</sub>)alkyl;

R<sub>4</sub> represents:  
30 hydrogen,  
methyl, or  
hydroxymethyl;

with the proviso that when R<sub>4</sub> is hydrogen or hydroxymethyl, then simultaneously R is methoxymethyl and R<sub>1</sub> is methyl;

and the pharmaceutically addition salts thereof.

This invention includes also a process for  
5 preparing the compounds of this invention from the  
corresponding starting compounds of formula (II)

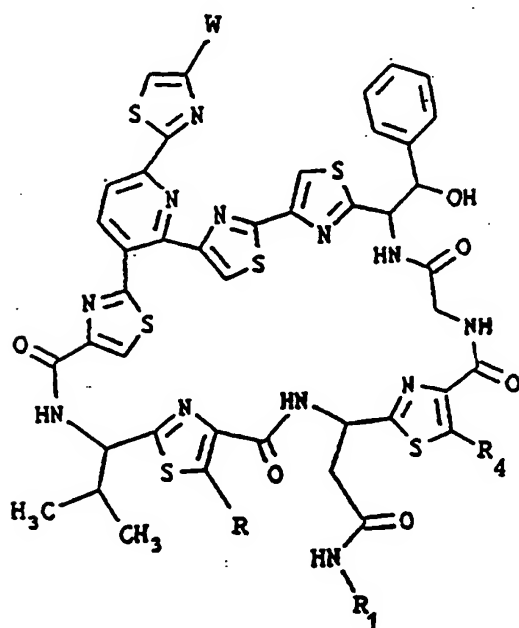
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II

wherein W is a carboxylic function or an activated ester thereof.

Antibiotic GE 2270 is prepared by culturing a sample of Planobispora rosea ATCC 53773 or a producing variant or mutant thereof and isolating the desired  
5 antibiotic substance from the mycelium and/or the fermentation broth. Planobispora rosea ATCC 53773 was isolated from a soil sample and deposited on June 14, 1988 with the American Type Culture Collection (ATCC), 12301 Parklawn Drive, Rockville, MD 20852 Maryland,  
10 U.S.A., under the provisions of the Budapest Treaty.

The strain has been accorded accession number ATCC 53773.

15 Antibiotic GE 2270 factor A is the main component of the antibiotic GE 2270 complex.

Antibiotic GE 2270 factor A and Planobispora rosea ATCC 53773 are described in European Patent  
20 Application Publication No. 359062.

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Recent studies showed that antibiotic GE 2270 factor A can be represented by the following general formula III



When antibiotic GE 2270 factor A is treated under selective hydrolysis conditions some derivatives named antibiotic GE 2270 factor A<sub>1</sub>, A<sub>2</sub> and A<sub>3</sub> are obtained. Said factors A<sub>1</sub>, A<sub>2</sub> and A<sub>3</sub> and the hydrolysis process for preparing them is disclosed in the European

Patent Application Publication No. 406745 and U.S.  
Patent Application No. 547,647.

5           Generally, the above mentioned hydrolytic  
conditions involve the use of mixtures of buffered or  
unbuffered aqueous acid media and polar organic  
solvents. The reaction temperature varies depending on  
factors such as the strength and the concentration of  
the acid employed, and is generally comprised between  
10       -10°C and 90°C. Also the reaction time varies  
considerably depending on parameters such as the  
temperature, the acid strength and its concentration;  
generally, it may vary from a few minutes to several  
15       hours.

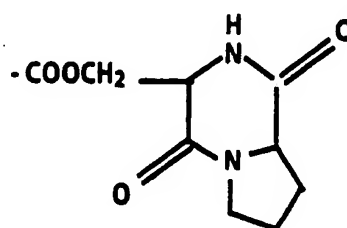
          In general, when milder hydrolysis conditions  
are employed, e.g. shorter reaction time and lower  
temperature or lower acid strength or concentration,  
20       antibiotic GE 2270 factor A<sub>1</sub> is normally obtained, while  
stronger hydrolysis conditions yield antibiotic GE 2270  
factor A<sub>2</sub>. To obtain antibiotic GE 2270 factor A<sub>3</sub>, still  
more drastic hydrolysis conditions are necessary.

25           While antibiotic GE 2270 factors A<sub>2</sub> and A<sub>3</sub> can  
be directly utilized as the starting materials for the  
production of the compounds of this invention,  
antibiotic GE 2270 factor A<sub>1</sub> is not suitable as the  
30       starting material for direct production of the compounds  
of this invention; however, it can be utilized as a  
precursor of the said starting materials as it will be  
explained further.

Antibiotic GE 2270 factors A<sub>2</sub> and factor A<sub>3</sub> are characterized by having an ester and a carboxy function respectively in the upper part of the molecule. In particular, it has been found that antibiotic GE 2270 factor A<sub>2</sub> and factor A<sub>3</sub> can be represented by the above defined formula II wherein:

W represents COOH (antibiotic GE 2270 factor A<sub>3</sub>) or the ester moiety (antibiotic GE 2270 factor A<sub>2</sub>)

15



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R is methoxymethyl,  
R<sub>1</sub> is methyl and  
R<sub>4</sub> is methyl.

Both antibiotic GE 2270 factor A<sub>2</sub> and factor A<sub>3</sub> (and mixture thereof) can be used as suitable starting materials for the production of the compounds of the invention, even if factor A<sub>3</sub> is the preferred one. Factor A<sub>2</sub> may be employed directly as an activated ester or may be converted to factor A<sub>3</sub> by drastic acid hydrolysis conditions, as mentioned above, or by basic

hydrolysis with diluted alkali (as described in European Patent Application Publication No. 406745 and U.S. Patent Application No. 547,647).

5           It was recently found (European Patent Application Publication No.451486 and U.S. Patent Application No. 665,612) that other minor components can be isolated from the cultures of Planobispora rosea ATCC 53773 or an antibiotic GE 2270 producing variant or  
10 mutant thereof. In particular, they are found in the mycelium and also in the fermentation broths of the cultured microorganism.

15           A preferred procedure for recovering said minor components of antibiotic GE 2270 from the mycelium includes extracting the filtered or centrifugated mycelium with a water-miscible organic solvent, concentrating the extracts and recovering the crude  
20 antibiotic substance by precipitation, optionally with the addition of a precipitating agent, by extraction of the aqueous residue with a water-immiscible organic solvent or by adsorption chromatography followed by elution of the desired product from the absorption  
25 matrix.

30           It was recently found (European Patent Application No. 91114667.8) that a further minor component (factor C<sub>2a</sub>) can be isolated from the same culture of Planobispora rosea ATCC 53773 described above.

The physico-chemical characteristics of antibiotic GE 2270 C<sub>2a</sub> are the following:



A) The ultraviolet absorption spectrum recorded with a Perkin Elmer Model 320 spectrometer exhibit the following absorption maxima:

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Solvent	UV max (nm)
0.1 M HCl	245-250 (shoulder) 300-315
0.1 M KOH	245-250 (shoulder) 300-315
Phosphate buffer pH 7.38	245-250 (shoulder) 300-315
Methanol	245-250 (shoulder) 300-315

25

30

B) The  $^1\text{H}$ -NMR spectrum of antibiotic GE 2270 factor  $\text{C}_{24}$  was recorded at 250 MHz with a Bruker spectrometer. The spectrum of the antibiotic in  $\text{DMSO-d}_6$  (hexadeuterodimethylsulfoxide) using TMS as the internal standard (0.00 ppm) exhibits the following groups of signals [ $\delta$ , ppm, m] (s=singlet, d=doublet, t=triplet, m=multiplet, Py=pyridine, Tz=thiazole)

9.03, d, (NH); 8.70, d, (2NH's); 8.60, s, 8.54, s, 8.29, s, and 7.38, s, (Tz CH's); 8.48, m, (glycine NH); 8.43, d, and 8.27, d, (Py CH's);

5 7.35-7.20, m, (aromatic CH's and primary amide NH); 6.98, s (primary amide NH); 6.04, d, (OH); 5.80, t (OH); 5.35-5.15, m, ( $\alpha$ CH's); 5.04, m, (phenylserine  $\beta$ CH); 4.98, s [ $\text{CH}_2(\text{OCH}_3)$ ]; 4.87, d, [ $\text{CH}_2(\text{OH})$ ]; 4.81, m and 4.56, m, (oxazoline  $\text{CH}_2$ ); 4.35-3.75, m, ( $\text{CH}_2$  of glycine and prolineamide CH's); 3.39, s, ( $\text{OCH}_3$ ); 2.71, m, and 1.30, m, ( $\text{CH}_2$  of asparagine); 2.48, d, ( $\text{NCH}_3$  of N-methylasparagine); 2.22-1.80, m, (isopropyl CH and prolineamide CH's); 0.88 and 0.84, d, (valine  $\text{CH}_3$ 's)

15 C) Antibiotic GE 2270 factor  $\text{C}_{2a}$  shows retention time ( $R_t$ ) of 12.6 min and retention time relative to antibiotic GE 2270 factor A ( $R_t$  16.6 min) of 0.76 when analyzed with the following reverse phase HPLC system:

20 Column: Bakerbond® C8 (5  $\mu\text{m}$ ) 4.6x250 mm (Bakerbond® is a trade name for reverse phase octylsilyl silica gel HPLC columns supplied by J.T. Baker Research Product, Phillipsburg, New Jersey 08865 USA)

25

Flow rate: 1.8 ml/min

30 Phase A:  $\text{CH}_3\text{CN}$ :tetrahydrofuran:40 mM  $\text{HCOONH}_4$   
40:40:20

Phase B:  $\text{CH}_3\text{CN}$ :tetrahydrofuran:40 mM  $\text{HCOONH}_4$   
10:10:80

Elution: linear gradient from 20% to 30% of Phase A in 20 min

Detection: UV 254 nm

5    D)    The main FAB-MS peak of antibiotic GE 2270  
factor C<sub>2a</sub> is 1306 daltons. This corresponds  
most likely to the lowest isotope of the  
protonated molecular ion. The analysis was  
10    performed on a Kratos MS-50 double focusing mass  
spectrometer, using 8 kV accelerating voltage  
and a saddle field atom gun with Xe gas ( $2 \times 10^{-5}$   
torr pressure indicated on the source ion gauge)  
at 6 kV voltage and 1 mA current. The antibiotic  
15    for the FAB-MS analysis was mixed with a  
thioglycerol matrix containing 0.1 M acetic  
acid.

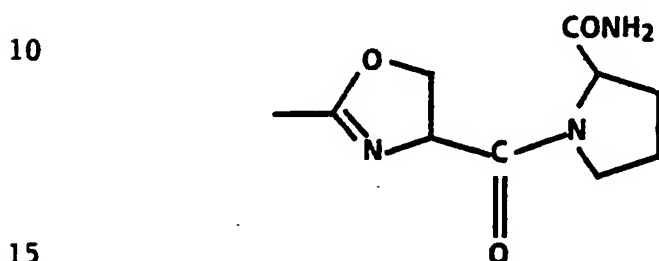
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Some of said minor components of antibiotic GE 2270 (i.e. factors B<sub>1</sub>, B<sub>2</sub>, C<sub>1</sub>, C<sub>2</sub>, C<sub>2a</sub>, D<sub>1</sub>, D<sub>2</sub> and E) may be represented by the general formula II mentioned above wherein

5 W represents the moiety:



R represents respectively hydrogen for GE 2270 factors C<sub>1</sub> and D<sub>1</sub>, methyl for factor B<sub>2</sub>, hydroxymethyl for factors D<sub>2</sub> and E and methoxymethyl for factors B<sub>1</sub>, C<sub>2</sub> and C<sub>2a</sub>;

20

R<sub>1</sub> represents hydrogen for GE 2270 factors B<sub>1</sub>, D<sub>1</sub> and E and methyl for GE 2270 factors B<sub>2</sub>, C<sub>1</sub>, C<sub>2</sub>, C<sub>2a</sub> and D<sub>2</sub>; and

25

R<sub>4</sub> represents hydrogen for GE 2270 factor C<sub>2</sub>, methyl for GE 2270 factors B<sub>1</sub>, B<sub>2</sub>, C<sub>1</sub>, D<sub>1</sub>, D<sub>2</sub> and E and hydroxymethyl for factor C<sub>2a</sub>.

30

When antibiotic GE 2270 factors D<sub>1</sub>, D<sub>2</sub> and E or mixture thereof are treated by the same hydrolytic process outlined above (and described in European Patent Application Publication No. 406745 and U.S. Patent Application No. 547,647) for preparing antibiotic

GE 2270 factors A<sub>2</sub> and A<sub>3</sub> from antibiotic GE 2270 factor A, the common moiety W cited above is hydrolyzed to a carboxy moiety leaving the substituents R, R<sub>1</sub> and R<sub>4</sub> unaltered.

5           Therefore, the derivatives of formula II wherein W is a carboxy or an activated ester function, R is hydrogen, hydroxymethyl or methoxymethyl, R<sub>1</sub> is hydrogen or methyl and R<sub>4</sub> is hydrogen, methyl or hydroxymethyl, with the proviso that when R<sub>4</sub> is hydrogen or  
10           hydroxymethyl then R is methoxymethyl and R<sub>1</sub> is methyl, can be used as starting material of the present invention. It has to be clear that as with other  
15           microorganisms, the characteristics of the GE 2270 producing strains are subject to variation. For example, artificial variants and mutants of the strain can be obtained by treatment with various known mutagens, such as U.V. rays, X-rays, high frequency waves, radioactive rays, and chemicals such as nitrous acid, N-methyl-N'-  
20           nitro-N-nitroso-guanidine, and many others. All natural and artificial variants and mutants which belong to a species of the genus Planobispora and produce antibiotic GE 2270 are deemed equivalent to strain Planobispora rosea ATCC 53773 for the purposes of this invention.  
25

          As used herein, the term "alkyl", either alone or in combination with other substituents, includes both straight and branched hydrocarbons groups; more  
30           particularly, "(C<sub>1</sub>-C<sub>14</sub>)alkyl" represents a straight or branched aliphatic hydrocarbon chain of 1 to 14 carbon atoms such as methyl, ethyl, propyl, 1-methylethyl, butyl, 1-methylpropyl, 1,1-dimethylethyl, pentyl, 1-methylbutyl, 2-methylbutyl, 1-hexyl, 2-hexyl, 3-hexyl, 3,3-dimethyl-1-butyl, 4-methyl-1-pentyl and 3-methyl-1-pentyl, heptyl, octyl, nonyl, decyl, undecyl,

dodecyl, tridecyl and tetradecyl; likewise, "(C<sub>1</sub>-C<sub>4</sub>)alkyl" represents a straight or branched hydrocarbon chain of 1 to 4 carbon atoms such as those alkyl of 1 to 4 carbons exemplified above.

5           As described above the "(C<sub>1</sub>-C<sub>14</sub>)alkyl" moiety may bear 1 to 3 substituents.

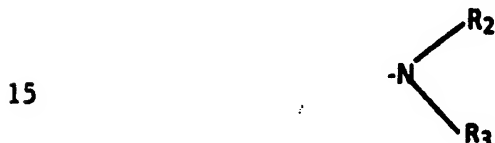
10           The term "halo" represents a halogen atom radical selected from fluoro, chloro, bromo and iodo.

15           As used herein, the term "(C<sub>3</sub>-C<sub>6</sub>)alkenyl" means an alkylene radical having three to six carbon atoms and a double bond; it comprises propenyl, 3-butenyl, 2-butenyl, 2-methylpropenyl, 2-pentenyl, 3-hexenyl and so on, which may be optionally substituted with a carboxy or a sulfo group.

20           The expression "a nitrogen containing 5-6 membered heterocyclic ring which may contain 1 to 3 further heteroatoms selected from N, S and O" according to the present invention includes unsaturated, partially saturated and wholly saturated ring systems such as pyridine, pyrimidine, pyrazine, pyrrolidine, piperidine, 25   piperazine, oxazole, oxazoline, oxazolidine, pyrazoline, pyrazolidine, thiazolidine, morpholine, thiomorpholine, pyrrole, pyrroline, imidazole, imidazolidine, thiadiazole, oxadiazole and tetrazole.

30           In said "nitrogen containing 5-6 membered heterocyclic ring" 1 to 3 ring carbons may optionally bear a group carboxy, sulfo, carboxy(C<sub>1</sub>-C<sub>4</sub>)alkyl and sulfo(C<sub>1</sub>-C<sub>4</sub>)alkyl and the ring nitrogen atom may optionally be substituted by (C<sub>1</sub>-C<sub>4</sub>)alkyl, carboxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, sulfo(C<sub>1</sub>-C<sub>4</sub>)alkyl, and benzyl.

The expression "fully saturated 5-7 membered nitrogen containing heterocyclic ring wherein the nitrogen atom may be optionally substituted by (C<sub>1</sub>-C<sub>4</sub>)alkyl or benzyl" identifies a fully saturated heterocycle of 5-7 members containing a nitrogen atom which can be optionally substituted by (C<sub>1</sub>-C<sub>4</sub>)alkyl or benzyl wherein the carbon skeleton may optionally bear one or two substituents selected from (C<sub>1</sub>-C<sub>4</sub>)alkyl, carboxy and sulfo. Said heterocyclic rings are connected with the nitrogen moiety of the rest



through a bond between the same nitrogen moiety and a carbon atom of the heterocyclic rest. Examples of said radicals are: 1-methyl-4-pyrrolidinyl, 3-piperidinyl, 1-ethyl-4-piperidinyl, 1-benzyl-2,6-dimethyl-4-piperidinyl, and 4-carboxy-1-methyl-2-piperidinyl.

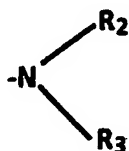
25 When R<sub>2</sub> and R<sub>3</sub> taken together with the adjacent nitrogen atom represent "a fully saturated 5-7 membered heterocyclic ring which may optionally contain a further heteroatom selected from O, S and N" this expression includes, for instance, the following heterocyclic groups: pyrrolidino, morpholino, piperidino, piperazino, thiomorpholino, pyrazolidino, 1,3-oxazolidino, 1,3-thiazolidino and hexahydroazepino. When the further heteroatom is N it may optionally bear a substituent selected from (C<sub>1</sub>-C<sub>4</sub>)alkyl, benzyl, carboxy, carboxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, sulfo and sulfo(C<sub>1</sub>-C<sub>4</sub>)alkyl.

The term "1-deoxy-1-glucityl" identifies a compound of formula (I) wherein Y is a radical deriving from glucamine, i.e. 1-amino-1-deoxy-glucitol. The term  
5 "2-deoxy-2-glucosyl" identifies a compound of formula (I) wherein Y is a radical deriving from glucosamine, i.e. 2-amino-2-deoxyglucose.

10 A preferred group of compounds of the invention is represented by those compounds of formula I wherein R represents methoxymethyl, R<sub>1</sub> and R<sub>4</sub> represent a methyl group and the other substituents are as defined above.

15 A further preferred group of compounds of the invention are those compounds of formula I wherein R represents methoxymethyl, R<sub>1</sub> and R<sub>4</sub> represent a methyl group, and Y represents a group of formula

20



25

wherein R<sub>2</sub> is hydrogen and R<sub>3</sub> is defined as above.

30 A further preferred group of compounds of the invention is represented by those compounds of formula I wherein R is methoxymethyl, R<sub>1</sub> and R<sub>4</sub> represent a methyl group and Y is an amino moiety which derive from a natural amino acid such as for example glycine, ornithine, serine, aspartic acid, tyrosine, leucine, phenylalanine, methionine, proline, threonine, lysine,

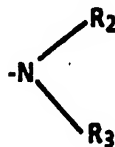


or a synthetic dipeptide such as glycyllysine, serylproline, glycyprolinamide, tyrosylprolinamide, threonylprolinamide, leucylprolinamide.

5 A further preferred group of compounds comprises those compounds of formula I wherein R is methoxymethyl, R<sub>1</sub> and R<sub>4</sub> are methyl, Y is a group NR<sub>2</sub>R<sub>3</sub> wherein R<sub>2</sub> is hydrogen and R<sub>3</sub> is a linear alkyl chain preferably of 3 to 12 carbons, more preferably of 3 to 7 carbons  
10 substituted with a group selected from COOH, SO<sub>3</sub>H and PO<sub>3</sub>H<sub>2</sub>.

The most preferred compound is represented by  
15 the formula I wherein R is methoxymethyl, R<sub>1</sub> and R<sub>4</sub> are methyl and Y is a group NR<sub>2</sub>R<sub>3</sub> wherein R<sub>2</sub> is hydrogen and R<sub>3</sub> is CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>-COOH.

20 A further preferred group of compounds of the invention are those compounds of formula I wherein R represents hydrogen, hydroxymethyl and methoxymethyl, R<sub>1</sub> represents hydrogen or a methyl group, and Y represents  
25 a group of formula



wherein R<sub>2</sub> is hydrogen and R<sub>3</sub> and R<sub>4</sub> are defined as above.

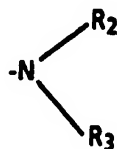
A further preferred group of compounds of the invention is represented by those compounds of formula I wherein R is hydrogen, hydroxymethyl or methoxymethyl, R<sub>1</sub> represents hydrogen or a methyl group, R<sub>4</sub> is hydrogen, methyl or hydroxymethyl with the proviso that when R<sub>4</sub> is hydrogen or hydroxymethyl then R is methoxymethyl and R<sub>1</sub> is methyl, and Y is an amino moiety which derive from a natural amino acid such as for example glycine, ornithine, serine, aspartic acid, tyrosine, leucine, phenylalanine, methionine, proline, threonine, lysine, or a synthetic dipeptide such as glycyllysine, serylproline, glycylprolinamide, tyrosylprolinamide, threonylprolinamide, leucylprolinamide.

A further preferred group of compounds comprises those compounds of formula I wherein R is hydrogen, hydroxymethyl or methoxymethyl, R<sub>1</sub> is hydrogen or methyl, R<sub>4</sub> is hydrogen, methyl or hydroxymethyl with the proviso that when R<sub>4</sub> is hydrogen or hydroxymethyl then R is methoxymethyl and R<sub>1</sub> is methyl, Y is a group NR<sub>2</sub>R<sub>3</sub> wherein R<sub>2</sub> is hydrogen and R<sub>3</sub> is a linear alkyl chain preferably of 3 to 12 carbons, more preferably of 3 to 7 carbons substituted with a group selected from COOH, SO<sub>3</sub>H and PO<sub>3</sub>H<sub>2</sub>.

The last preferred group of compounds is represented by the formula I wherein R hydrogen, hydroxymethyl or methoxymethyl, R<sub>1</sub> is hydrogen or methyl, R<sub>4</sub> is as defined above and Y is a group NR<sub>2</sub>R<sub>3</sub> wherein R<sub>2</sub> is hydrogen and R<sub>3</sub> is CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>-COOH.

Representative examples of the compounds of the invention, include those compounds of formula I wherein R, R<sub>1</sub>, R<sub>4</sub> and Y are as defined above and

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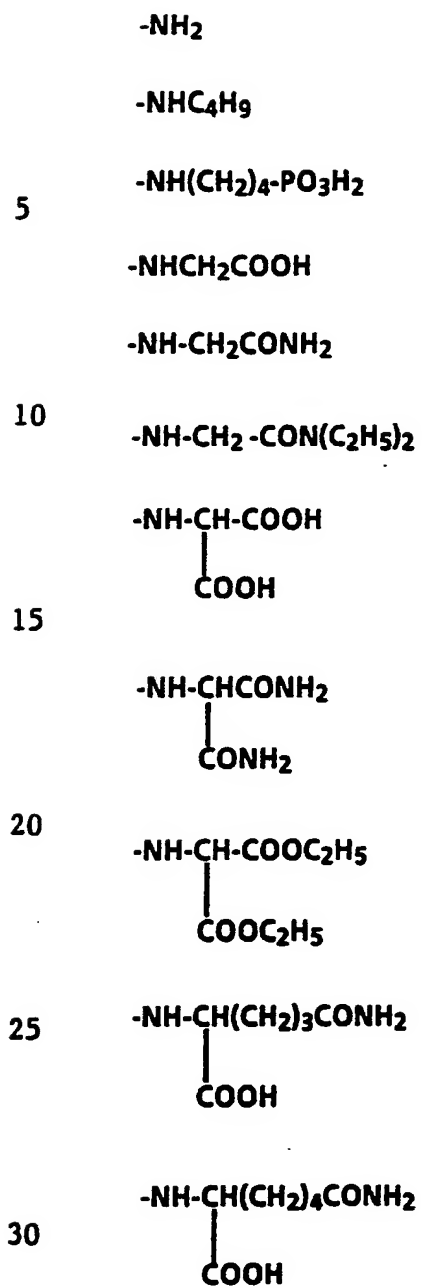
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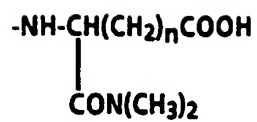
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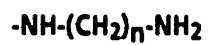
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5            wherein n is 2, 3 or 4



10             $\text{-NH-(CH}_2\text{)}_n\text{-NHCH}_3$

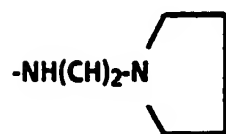


15             $\text{-NH-(CH}_2\text{)}_n\text{-N(C}_2\text{H}_5\text{)}_2$

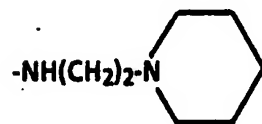
20             $\text{-HN-(CH}_2\text{)}_n\text{-N(CH}_3\text{)}(\text{C}_2\text{H}_5)$

             wherein n is 2, 3, 4, 5, 6, 7 or 8

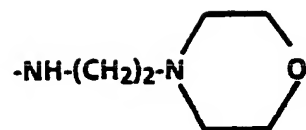
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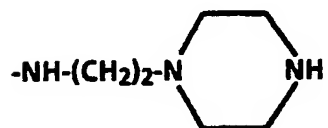
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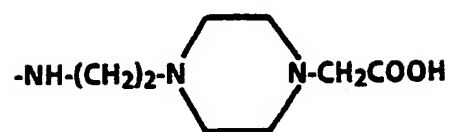
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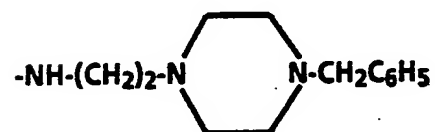
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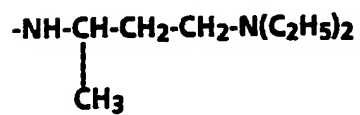
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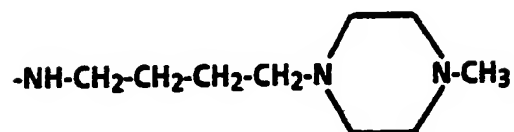
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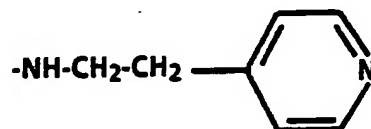
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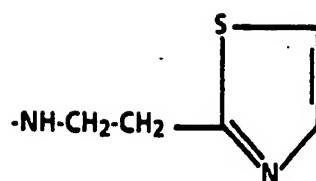
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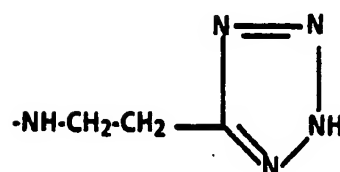
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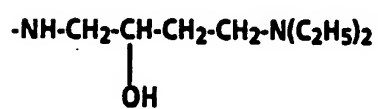
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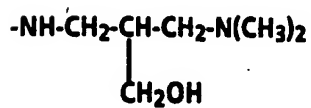
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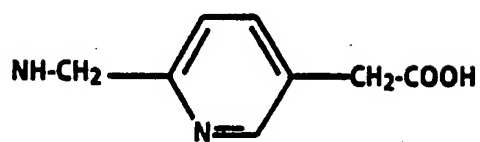
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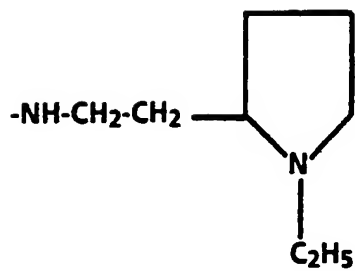


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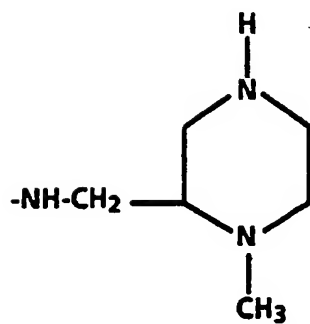
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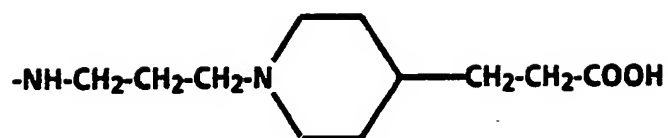
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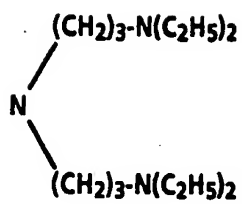
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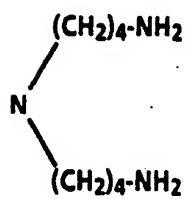
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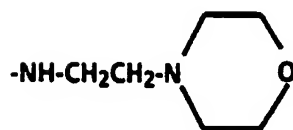


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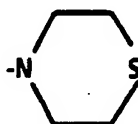
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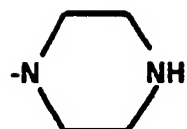


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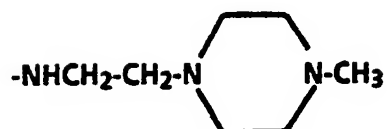


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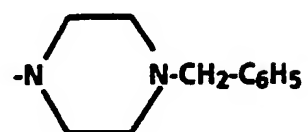
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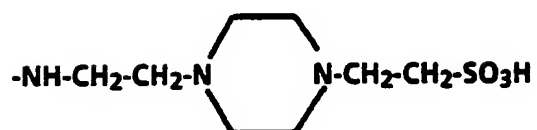
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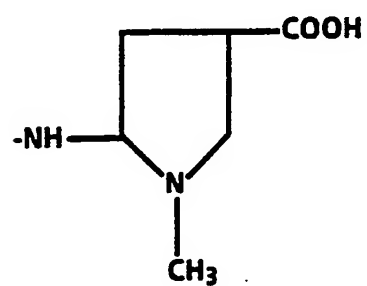
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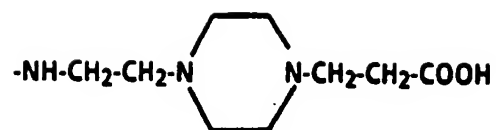


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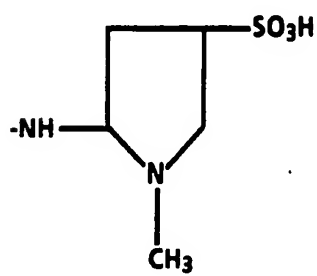


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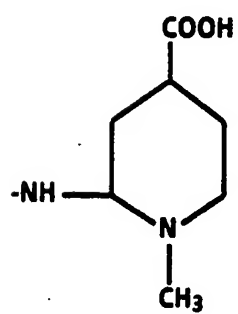


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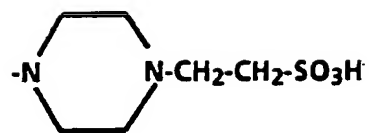
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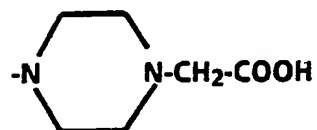
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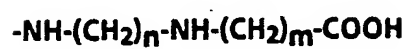
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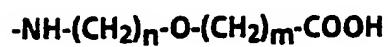
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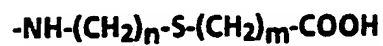
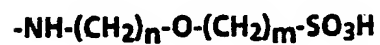
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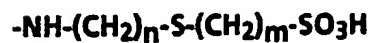
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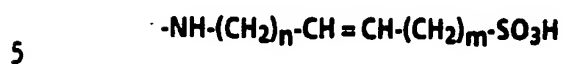
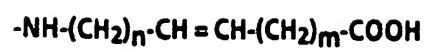


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wherein n is 2, 3, 4 or 5 and m is 1, 2, 3 or 4

30



wherein n is 1, 2 or 3 and m is 0, 1 or 2

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The compounds of the invention can form salts according to conventional procedures.

In particular, those compounds of formula I  
5 wherein the group  $-NR_2R_3$  contains further amine functions form acid addition salts.

In addition, those compounds of the invention  
10 which contain acid functions in the  $-NR_2R_3$  moiety may also form base addition salts.

In general, those compounds of the invention which contain acid and basic functions can form internal salts. For the scope of the present invention the  
15 "internal salts" are encompassed by the definition of the "non-salt" form.

Preferred addition salts of the compounds of this invention are the pharmaceutically acceptable acid and/or base addition salts.

20 With the term "pharmaceutically acceptable acid and/or base addition salts" are intended those salts with acids and/or bases which from biological, manufacturing and formulation standpoint are compatible with the pharmaceutical practice as well as with the use  
25 in the animal growth promotion.

Representative and suitable acid addition salts of the compounds of formula I include those salts formed by standard reaction with both organic and inorganic  
30 acids such as, for example, hydrochloric, hydrobromic, sulfuric, phosphoric, acetic, trifluoroacetic, trichloroacetic, succinic, citric, ascorbic, lactic, maleic, fumaric, palmitic, cholic, pantoic, mucic, glutamic, camphoric, glutaric, glycolic, phthalic, tartaric, lauric, stearic, salicylic, methanesulfonic,

dodecylsulfonic acid (estolic acid), benzenesulfonic, sorbic, picric, benzoic, cinnamic and the like acids.

Representative examples of these bases are:  
alkali metal or alkaline-earth metal hydroxide such  
5 sodium, potassium, and calcium hydroxide; ammonia and  
organic aliphatic, alicyclic or aromatic amines such as  
methylamine, dimethylamine, trimethylamine,  
2-amino-2-hydroxymethyl-1,3-propanediol (TRIS), picoline  
and basic aminoacids such as lysine, ornithine, arginine  
10 and histidine.

The transformation of the free amino or non-salt  
compounds of the invention into the corresponding  
addition salts, and the reverse, i.e. the transformation  
15 of an addition salt of a compound of the invention into  
the non-salt or free amino form, are within the ordinary  
technical skill and are encompassed by the present  
invention.

For instance, a compound of formula I can be  
20 transformed into the corresponding acid or base  
addition-salt by dissolving the non-salt form in an  
aqueous solvent and adding a slight molar excess of the  
selected acid or base. The resulting solution or  
suspension is then lyophilized to recover the desired  
25 salt. Instead of lyophilizing, in some instances, it is  
possible to recover the final salt by extraction with an  
organic solvent, concentration to a small volume of the  
separated organic phase and precipitation by adding a  
30 non-solvent.

In case the final salt is insoluble in an  
organic solvent where the non-salt form is soluble it is  
recovered by filtration from the organic solution of the  
non-salt form after addition of the stoichiometric  
amount or a slight molar excess of the selected acid or  
base.

The non-salt form can be prepared from a corresponding acid or base salt dissolved in an aqueous solvent which is then neutralized to free the non-salt form. This is then recovered for instance by extraction  
5 with an organic solvent or is transformed into another base or acid addition salt by adding the selected acid or base and working up as above.

When following the neutralization desalting is necessary, a common desalting procedure may be employed.  
10

For example, column chromatography on controlled pore polydextrane resins (such as Sephadex LH 20) or silanized silica gel may be conveniently used. After eluting the undesired salts with an aqueous solution,  
15 the desired product is eluted by means of linear gradient or step-gradient of a mixture of water and a polar or apolar organic solvent, such as acetonitrile/water from 50:50 to about 100% acetonitrile.

20

As is known in the art, the salt formation either with pharmaceutically acceptable acids (bases) or non-pharmaceutically acceptable acids (bases) may be used as a convenient purification technique. After  
25 formation and isolation, salt form of a compound of formula I can be transformed into the corresponding non-salt or into a pharmaceutically acceptable salt.

30 In some instances the acid addition salt of a compound of formula I is more soluble in water and hydrophilic solvents and has an increased chemical stability.

However, in view of the similarity of the properties of the compounds of formula I and their



salts, what is said in the present application when dealing with the biological activities of the compounds of formula I applies also to their pharmaceutically acceptable salts, and viceversa.

5

In view of their properties, the compounds of the invention can be used as active ingredients in the preparation of medicaments for human or animal treatment.

10

In particular, the amide derivatives of the antibiotic GE 2270 compounds of formula I are antimicrobial agents mainly active against gram positive bacteria and gram positive as well as gram negative anaerobes.

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A general procedure for preparing a compound of  
this invention is represented by the reaction  
(amidation) of a suitable antibiotic GE 2270 compound  
5 having formula (II)

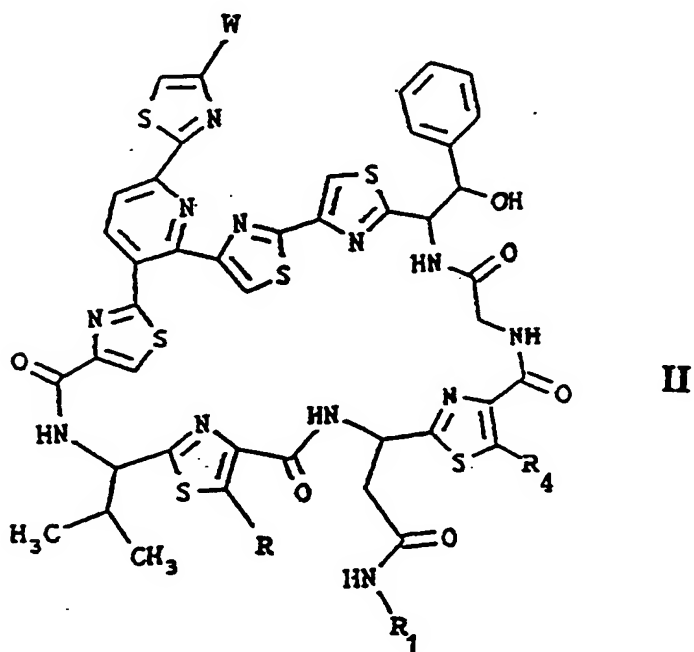
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wherein

- W represents a carboxy or an activated ester function;
- R represents hydrogen, hydroxymethyl or methoxymethyl;
- 5 R<sub>1</sub> represents hydrogen or methyl;
- R<sub>4</sub> represents hydrogen, methyl or hydroxymethyl,

with the proviso that, when R<sub>4</sub> represents hydrogen or hydroxymethyl, then simultaneously R is methoxymethyl and R<sub>1</sub> is methyl; with a selected amine of formula  
10  $\text{HNR}_2\text{R}_3$  wherein R<sub>2</sub> and R<sub>3</sub> have the same meanings as above in an inert organic solvent and, when W is carboxy, in the presence of a condensing agent.

15 In carrying out the amidation for preparing the compounds of this invention, sometimes, it is convenient to protect the functions of the reactants which are not involved in the amidation reaction but could result  
20 sensitive to the reaction conditions or negatively affect the reaction course, for instance, yielding undesired side-product.

25 Furthermore, when the amino acid contains further reactive functions such as amino, carboxy or mercapto groups which may interfere with the course of the amidation, these are protected by means of methods known per se in the art such as those described in  
30 references books like E. Gross and J. Meienhofer "The Peptides", Vol. 3, Academic Press, New York, 1981 and M. Bodanszky and A. Bodanszky "The Practice of Peptide Synthesis, Springer-Verlag, Berlin Heidelberg, 1984. These protecting groups must be stable at the conditions the amidation reaction takes place and must be easily removable at the end of the reaction without affecting

either the newly formed amide bond or any other part of the molecule.

Representative examples of N-protecting groups which may be advantageously used in the process of the invention for protecting an amino function are carbamate forming reagents characterized by the following oxycarbonyl groups: 1,1-dimethylpropynyl-oxycarbonyl, t-butyloxycarbonyl, vinyloxycarbonyl, aryloxycarbonyl, cinnamyloxycarbonyl, benzyloxycarbonyl, p-nitrobenzyloxycarbonyl-3,4-dimethoxy-6-nitrobenzyloxycarbonyl, 2,4-dichlorobenzyloxycarbonyl, 5-benzisoxazolylmethyloxycarbonyl, 9-anthranylmethyloxycarbonyl, diphenylmethyloxycarbonyl, isonicotinylloxycarbonyl, diphenylmethyloxycarbonyl, isonicotinylloxycarbonyl, S-benzyloxycarbonyl, and the like.

A suitable protection for reactive carboxylic acid function is, for instance, by forming an ester function.

The man skilled in the art is capable, also on the basis of the present disclosure, of deciding which functions of the amine  $\text{HNR}_2\text{R}_3$  need to be protected, how they must be protected and the proper deprotection reaction which is necessary to free the final compound.

As it is appreciated by the skilled technician, the ultimate choice of the specific protecting group depends on the characteristics of the particular amide derivative which is desired. In fact, this amide function of the final compound should be stable at the condition of removal of the protecting group(s).

Since the conditions of removal of the different protecting groups are known, the skilled technician is capable of selecting the proper protecting group.

5           Inert organic solvents useful for the condensation reaction are those solvents which do not unfavorably interfere with the reaction course and are capable of at least partially solubilizing the antibiotic starting material.

10           Examples of said inert solvents are organic amides, ethers of glycols and polyols, phosphoramides, sulfoxides. Preferred examples of inert solvents are: dimethylformamide, dimethoxyethane,  
15           hexamethylphosphoramide, dimethylsulfoxide, dioxane, and mixtures thereof.

Sometimes, water is compatible with the reaction conditions.

20           The condensing agent in the process of the invention when W is carboxy is one suitable for forming amide bonds in organic compounds and in particular in peptide synthesis.

25           Representative and preferred examples of condensing agents are (C<sub>1</sub>-C<sub>4</sub>)alkyl, phenyl or heterocyclic phosphorazidates such as, diphenylphosphorazidate (DPPA), diethyl-phosphorazidate, di(4-nitrophenyl)phosphorazidate, dimorpholylphosphorazidate and diphenylphosphorochloridate or  
30           benzotriazol-1-yl-oxy-tripyrrolidinophosphoniumhexafluorophosphate (PyBOP). The preferred condensing agent is diphenyl phosphorazidate (DPPA).

In the process of the invention, the amine reactant  $\text{HNR}_2\text{R}_3$  is normally used in a slight molar excess.

In general, a 1- to 2-fold molar excess is used  
5 while a 1.2- to 1.5-fold molar excess is preferred.

For the amidation to proceed, it is necessary that the amine  $\text{HNR}_2\text{R}_3$  be capable of forming a salt with the carboxy function of the antibiotic starting  
10 material. In case the amine  $\text{HNR}_2\text{R}_3$  is not strong enough to form such a salt in the selected reaction medium, it is necessary to add a salt-forming base to the reaction mixture at least in an equimolecular amount with the  
15 antibiotic starting material.

Examples of said salt-forming bases are tertiary organic aliphatic or alicyclic amines such as trimethylamine, triethylamine, N-methyl pyrrolidine or heterocyclic bases such as picoline, and the like.

20 The condensing agent is generally employed in a slight molar excess such as from 1.1 to 1.5 and preferably is 1.2 times the antibiotic GE 2270 starting compound.

In addition, the amine reactant  $\text{HNR}_2\text{R}_3$  may also  
25 conveniently be introduced in the reaction medium as a corresponding acid addition salt, e.g. the hydrochloride. In this case, at least a double molar proportion and preferably a 2 to 3 fold molar excess of  
30 a strong base capable of freeing the  $\text{HNR}_2\text{R}_3$  amine from its salts, is used. Also in this case, the suitable base is a tertiary organic aliphatic or alicyclic amine like those exemplified above. In fact, at least in some instances, the use of salt of the amine  $\text{HNR}_2\text{R}_3$ , which is then freed in situ with the above mentioned bases, is

greatly preferred especially when the salt is more stable than the corresponding free amine.

5 The reaction temperature will vary considerably depending on the specific starting materials and reaction conditions. In general, it is preferred to conduct the reaction at temperatures between 0-20°C.

Also the reaction time vary considerably depending on the other reaction parameters. In general  
10 the condensation reaction is completed in about 5-24 h.

In any case, the reaction course is monitored by TLC or preferably by HPLC according to methods known in the art.

15 On the basis of the results of these assays a man skilled in the art will be able to evaluate the reaction course and decide when to stop the reaction and start working up the reaction mass according to known per se techniques which include, for instance,  
20 extraction with solvents, precipitation by addition of non-solvents, etc., in conjunction with further separations and purifications by column chromatography.

As already said, when protection of the  $\text{HNR}_2\text{R}_3$  reactant is necessary, the protected final compound is  
25 then de-protected according to procedures which are known per se and mainly depends on the protecting group involved.

30 When an activated ester is used as the GE 2270 starting material, said ester is one wherein the esterified alcohol is providing a leaving group which can be readily displaced and substituted by the amine  $\text{HNR}_2\text{R}_3$  under reaction conditions which do not modify the other portions of the molecule. The amine reactant is

usually employed in a molar excess over the activated ester in a solvent which is selected from those mentioned above and the lower alkanols. The reaction temperature generally ranges between 0°C and 100°C.

5 Examples of the activated ester include lower alkyl esters wherein the lower alkyl moiety is optionally substituted by cyano and nitro, phenyl esters substituted by halo and nitro groups as well as the ester moiety contained in GE 2270 factor A<sub>2</sub>.

10

It is evident that in many instances a compound of the invention may be prepared in more than one way and that a compound of the invention may be transformed  
15 into another by means of known per se reactions.

For instance when the HNR<sub>2</sub>R<sub>3</sub> amine contains a carboxy or an ester function which can be further converted into the corresponding amide derivative, a  
20 desired compound of formula I may be prepared by condensing first said amine with the selected GE 2270 starting material and then converting the carboxy or ester function to amide by reaction with the appropriate  
25 amine.

25

The following tables list the structure formulas  
30 of some representative compounds of the invention (TABLE I) and their methods of preparation, (described in details in the Experimental Section), starting materials and reaction yields (TABLE II).



In the following Table (TABLE I) the structure formulas of representative examples of compounds of the invention are reported.

TABLE I


COMPOUND OF EXAMPLE NO.	Y	R	R <sub>1</sub>	R <sub>4</sub>
1	-NH CH <sub>2</sub> COOH	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
2	-NH CHCOOH   CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NH <sub>2</sub>	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
3	-NH CHCOOH   CH <sub>2</sub> OH	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
4	-NH CHCOOH   CH <sub>2</sub> COOH	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
5	-NH CH-COOH   CH <sub>2</sub> - 	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>

TABLE I (continued)

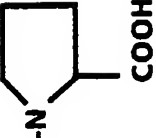
COMPOUND OF EXAMPLE NO.	Y	R	R <sub>1</sub>	R <sub>4</sub>
6	$\begin{array}{c} \text{-NH CHCOOH} \\   \\ \text{CH}_2\text{CH}(\text{CH}_3)_2 \end{array}$	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
7	$\begin{array}{c} \text{-NH CH-COOH} \\   \\ \text{CH}_2-\text{C}_6\text{H}_5 \end{array}$	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
8	$\begin{array}{c} \text{-NH CHCOOH} \\   \\ \text{CH}_2\text{CH}_2\text{SCH}_3 \end{array}$	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
9	$\begin{array}{c} \text{COOH} \\   \\ \text{N} \end{array}$ 	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
10	$\begin{array}{c} \text{-NH CHCOOH} \\   \\ \text{HO}-\text{CH}-\text{CH}_3 \end{array}$	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>

TABLE I (continued)

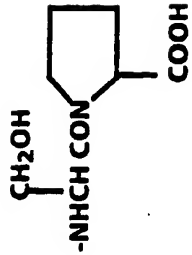
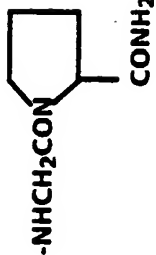
COMPOUND OF EXAMPLE NO.	Y	R	R <sub>1</sub>	R <sub>4</sub>
11	$\text{-NHCH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}(\text{NH}_2)\text{COOH}$	$\text{CH}_2\text{O-CH}_3$	$\text{CH}_3$	$\text{CH}_3$
12	$\text{-NHCH}_2\text{CONHCH}(\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{NH}_2)\text{COOH}$	$\text{CH}_2\text{O-CH}_3$	$\text{CH}_3$	$\text{CH}_3$
13		$\text{CH}_2\text{O-CH}_3$	$\text{CH}_3$	$\text{CH}_3$
14		$\text{CH}_2\text{O-CH}_3$	$\text{CH}_3$	$\text{CH}_3$

TABLE I (continued)

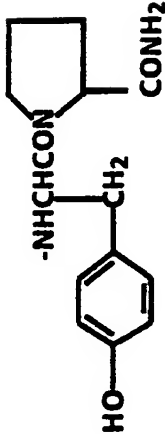
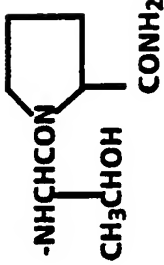
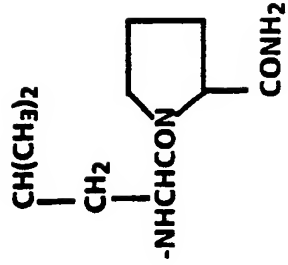
COMPOUND OF EXAMPLE NO.	Y	R	R <sub>1</sub>	R <sub>4</sub>
15		CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
16		CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
17		CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>

TABLE I (continued)

COMPOUND OF EXAMPLE NO.	Y	R	R <sub>1</sub>	R <sub>4</sub>
18	-NH CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOH	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
19	-NH CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOH	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
20	-NH CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOH	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
21	-NH CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOH	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
22	-NH CH <sub>2</sub> CH <sub>2</sub> SO <sub>3</sub> H	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
23	-NH CH <sub>2</sub> CH <sub>2</sub> SO <sub>3</sub> H	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>

TABLE I (continued)




COMPOUND OF EXAMPLE NO.	Y	R	R <sub>1</sub>	R <sub>4</sub>
24	-NH CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> PO <sub>3</sub> H <sub>2</sub>	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
25	-NH CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> PO <sub>3</sub> H <sub>2</sub>	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
26		CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
27		CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
28		CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>

TABLE I (continued)

COMPOUND OF EXAMPLE NO.	Y	R	R <sub>1</sub>	R <sub>4</sub>
29	$  \begin{array}{c}  \text{N-CH}_2\text{CH} \\    \\  \text{CH}_3  \end{array}  \begin{array}{c}  \text{CH} \\    \\  \text{OH}  \end{array}  \begin{array}{c}  \text{CH} \\    \\  \text{OH}  \end{array}  \begin{array}{c}  \text{CH} \\    \\  \text{OH}  \end{array}  \begin{array}{c}  \text{CH} \\    \\  \text{OH}  \end{array}  \begin{array}{c}  \text{CH}_2\text{OH}  \end{array}  $	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
30		CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
31	$  \begin{array}{c}  \text{NHCH}_2\text{CH}_2\text{CH}_2\text{NH} \\    \\  \text{C=O}  \end{array}  $	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
32	$  \begin{array}{c}  \text{NH} \\    \\  \text{CH}_2\text{CH}_2\text{N}(\text{CH}_3)_2  \end{array}  $	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>

TABLE I (continued)



COMPOUND OF EXAMPLE NO.	Y	R	R <sub>1</sub>	R <sub>4</sub>
33		CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
34		CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
35	-NH <sub>2</sub>	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
36	-NH CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NH <sub>2</sub>	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>



TABLE I (continued)

COMPOUND OF EXAMPLE NO.	Y	R	R <sub>1</sub>	R <sub>4</sub>
37	-NH CH <sub>2</sub> CHO	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
38	-NH CH <sub>2</sub> CH <sub>2</sub> NHCH <sub>2</sub> CH <sub>2</sub> COOH	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
39	-NH CH <sub>2</sub> CH <sub>2</sub> SCH <sub>2</sub> CH <sub>2</sub> COOH	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
40	-NH CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH = CHCOOH	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>
41	-NH CH <sub>2</sub> CH <sub>2</sub> OCH <sub>2</sub> CH <sub>2</sub> COOH	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>

TABLE I (continued)

COMPOUND OF EXAMPLE NO.	Y	R	R <sub>1</sub>	R <sub>4</sub>
42	-NH CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOH	CH <sub>2</sub> O-CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>2</sub> OH
43	-NH CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOH	H	H	CH <sub>3</sub>
44	-NH CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOH	CH <sub>2</sub> OH	CH <sub>3</sub>	CH <sub>3</sub>

Compounds No. 2, 11, 12, 34, 36 were isolated as trifluoroacetate salts

TABLE II



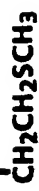

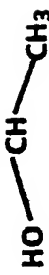
COMPOUND OF EXAMPLE NO.	STARTING MATERIALS (GE 2270 FACTOR + AMINE REACTANT)	METHOD	OVERALL YIELD
1	A <sub>3</sub> + HCl.NH <sub>2</sub> CH <sub>2</sub> COOEt	A <sub>1</sub>	80%
2	A <sub>3</sub> + HCl.NH <sub>2</sub> CHCOOMe   CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NH.Cbz	A <sub>1</sub>	72%
3	A <sub>3</sub> + HCl.NH <sub>2</sub> CHCOOMe   CH <sub>2</sub> OH	A <sub>1</sub>	70%
4	A <sub>3</sub> + HCl.NH <sub>2</sub> CHCOOMe   CH <sub>2</sub> COOMe	A <sub>1</sub>	54%
5	A <sub>3</sub> + HCl.NH <sub>2</sub> CH-COOMe   CH <sub>2</sub> - 	A <sub>1</sub>	70%
6	A <sub>3</sub> + HCl.NH <sub>2</sub> CHCOOMe   CH <sub>2</sub> CH(CH <sub>3</sub> ) <sub>2</sub>	A <sub>1</sub>	70%

TABLE II (continued)

COMPOUND OF EXAMPLE NO.	STARTING MATERIALS (GE 2270 FACTOR + AMINE REACTANT)	METHOD	OVERALL YIELD
7	A <sub>3</sub> + HCl.NH <sub>2</sub> CHCOOMe 	A <sub>1</sub>	60%
8	A <sub>3</sub> + HCl.NH <sub>2</sub> CHCOOMe 	A <sub>1</sub>	70%
9	A <sub>3</sub> + HCl.HN 	A <sub>1</sub>	75%
10	A <sub>3</sub> + HCl.NH <sub>2</sub> CHCOOMe 	A <sub>1</sub>	70%

**TABLE II (continued)**

TABLE II (continued)


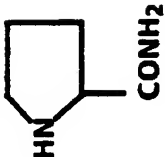
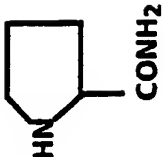
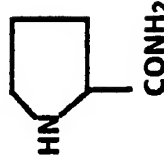
COMPOUND OF EXAMPLE NO.	STARTING MATERIALS (GE 2270 FACTOR + AMINE REACTANT)	METHOD	OVERALL YIELD
15	$A_3 + HCl.NH_2CHCON$ 	A	60%
	or $\underline{5} +$ 	C	70%
16	$\underline{10} +$ 	C	60%
17	$\underline{6} +$ 	C	65%

TABLE II (continued)

COMPOUND OF EXAMPLE NO.	STARTING MATERIALS (GE 2270 FACTOR + AMINE REACTANT)	METHOD	OVERALL YIELD
18	A <sub>3</sub> + HCl.NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOMe	A <sub>1</sub>	73%
19	A <sub>3</sub> + HCl.NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOMe	A <sub>1</sub>	77%
	A <sub>3</sub> + NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOH	B	70%
20	A <sub>3</sub> + PTSA.NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOMe	A <sub>1</sub>	75%
21	A <sub>3</sub> + PTSA.NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOMe	A <sub>1</sub>	70%
22	A <sub>3</sub> + NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>3</sub> H	B	20%

TABLE II (continued)


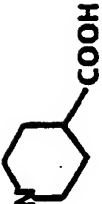
COMPOUND OF EXAMPLE NO.	STARTING MATERIALS (GE 2270 FACTOR + AMINE REACTANT)	METHOD	OVERALL YIELD
23	A <sub>3</sub> + NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>3</sub> H	B	25%
24	A <sub>3</sub> + NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> PO <sub>3</sub> H <sub>2</sub>	B	40%
25	A <sub>3</sub> + NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> PO <sub>3</sub> H <sub>2</sub>	B	35%
26	A <sub>3</sub> + NH <sub>2</sub> CH <sub>2</sub> - 	B	60%
27	A <sub>3</sub> + HN- 	B	50%



TABLE II (continued)

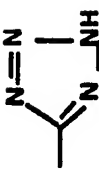
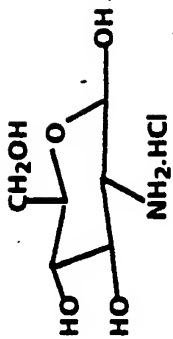
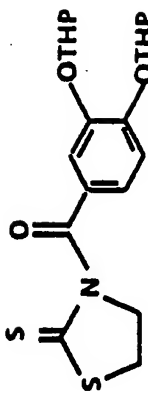
COMPOUND OF EXAMPLE NO.	STARTING MATERIALS (GE 2270 FACTOR + AMINE REACTANT)	METHOD	OVERALL YIELD
28	$A_3 + NH_2CH_2CH_2CH_2CH_2CH_2 -$ 	B	65%
29	$A_3 + NH - CH_2 - CH(CH_3) - CH(OH) - CH(OH) - CH_2OH$	A	20%
30	$A_3 +$ 	A	80%
31	$36 +$ 	C1	80%

TABLE II (continued)

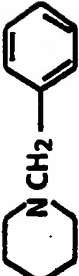

COMPOUND OF EXAMPLE NO.	STARTING MATERIALS (GE 2270 FACTOR + AMINE REACTANT)	METHOD	OVERALL YIELD
32	A <sub>3</sub> + NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> N(CH <sub>3</sub> ) <sub>2</sub>	A	75%
33	A <sub>3</sub> + NH <sub>2</sub> — 	A	60%
34	A <sub>3</sub> + NH— 	A <sub>1</sub>	50%
35	A <sub>3</sub> + NH <sub>3</sub> in MeOH	D	83%
36	A <sub>3</sub> + NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NH.Boc	A <sub>1</sub>	70%

TABLE II (continued)

COMPOUND OF EXAMPLE NO.	STARTING MATERIALS (GE 2270 FACTOR + AMINE REACTANT)	METHOD	OVERALL YIELD
37	A <sub>3</sub> + NH <sub>2</sub> CH <sub>2</sub> CH <div style="display: inline-block; vertical-align: middle; text-align: center;"><math>\begin{array}{c} \text{OCH}_3 \\ \diagup \\ \text{CH} \\ \diagdown \\ \text{OCH}_3 \end{array}</math></div>	A <sub>1</sub>	65%
38	37 + HCl.NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOCH <sub>2</sub> CH <sub>3</sub>	C <sub>1</sub>	20%
39	A <sub>3</sub> + TFA.NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SCH <sub>2</sub> CH <sub>2</sub> COOCH <sub>3</sub>	A <sub>1</sub>	33%
40	A <sub>3</sub> + TFA.NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH=CH-COOH	B	51%

TABLE II (continued)

COMPOUND OF EXAMPLE NO.	STARTING MATERIALS (GE 2270 FACTOR + AMINE REACTANT)	METHOD	OVERALL YIELD
41	A <sub>3</sub> + TFA.NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OCH <sub>2</sub> CH <sub>2</sub> COOH	B	37%
42	C <sub>2a</sub> + HCl.NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOCH <sub>3</sub>	F	40%
	NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOH	G	35%
43	D <sub>1</sub> + HCl.NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOCH <sub>3</sub>	H	50%
	NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOH	I	40%
44	D <sub>2</sub> + HCl.NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOCH <sub>3</sub>	J	35%
	NH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOH	K	30%

TFA = trifluoroacetic acid  
PTSA = p-toluenesulfonic acid

### HPLC Analysis

The following table (TABLE III) reports the  $R_t$  of representative examples of compounds of this invention.

Analysis were run with a Varian model 5000 LC pump equipped with a 10  $\mu$ l loop injector and a Varian 2050 variable wavelength detector at 254 nm.

Columns: Pre-column LiChroCart-LiChrosorb RP-8 (5  $\mu$ m) followed by a column LiChroCart 125-4 LiChrospher 100 RP-8 (5  $\mu$ m)

Eluents: A 0.05 M aq.  $\text{HCOONH}_4$   
B  $\text{CH}_3\text{CN}$   
C THF

Method A: isocratic 44% of B in A  
Flow rate: 0.7 ml/min

Method B: isocratic 40% of B in A  
Flow rate 0.7 ml/min

Method C: isocratic 38% of B in A  
Flow rate: 0.5 ml/min

Method D: isocratic 30% of B in A  
Flow rate: 0.7 ml/min

Method E: isocratic 38% of B in A  
Flow rate: 0.7 ml/min

**Method F:** gradient from 38 to 55% of B in A in  
11 min according to the following program

	Time (min)	%B in A
5	0	38
	6	38
	7	45
	10	45
	11	55
10		

Flow rate: 0.7 ml/min

**Method G:** gradient from 38 to 55% of B in A in 25  
min according to the following program

	Time (min)	%B in A
20	0	38
	6	38
	10	44
	15	44
	25	55

Flow rate: 0.7 ml/min

**Method H:** isochratic 55% of B in A  
Flow rate: 0.7 ml/min

**Method I:** isochratic 60% of B in A  
Flow rate: 0.7 ml/min

**Method L:** isochratic 48% of B in A  
Flow rate: 0.7 ml/min

Method M: gradient according to the following program:

	Time (min)	%A	%B	%C
5	0	74	10	16
	20	62	19	19

Flow rate: 0.7 ml/min

10

15

20

25

30

**TABLE III**  
**HPLC Analysis**

<b>Compound No.</b>	<b>Method</b>	<b>R<sub>t</sub> (min)</b>	<b>K</b>
1	A	2.56	0.92
2	B	4.09	1.15
3	C	6.21	1.12
4	D	14.70	1.79
5	E	5.99	1.28
6	A	4.05	1.46
7	A	4.52	1.63
8	A	3.44	1.24
9	E	5.32	1.18
10	E	4.22	0.90
11	F	14.30	3.05
12	F	5.99	1.28
13	E	4.88	1.04
14	G	14.09	3.01
15	G	17.60	3.76
16	G	13.77	2.94
17	G	23.75	5.07
18	G	7.49	1.60

**K = Relative Retention time**



TABLE III (continued)

## HPLC Analysis

Compound No.	Method	R <sub>t</sub> (min)	K
19	F	8.84	1.89
20	G	17.77	3.78
21	G	31.10	6.64
22	F	5.01	1.07
23	F	4.40	0.94
24	F	6.27	1.34
25	F	11.17	2.39
26	F	29.04	6.02
27	F	6.28	1.34
28	F	12.14	2.59
29	F	8.02	1.71
30	E	6.81-7.61 anomeric mixture	1.45-1.62 anomeric mixture
31	F	17.64	3.76
32	B	9.64	2.70
33	H	15.10	7.40
34	I	7.22	3.92
35	L	10.28	4.11
36	F	19.32	4.13

K = Relative Retention time

**TABLE III (continued)**  
**HPLC Analysis**

<b>Compound No.</b>	<b>Method</b>	<b>R<sub>t</sub> (min)</b>	<b>K</b>
37	F	14.56	3.11
38	F	11.00	2.35
39	F	9.48	2.02
40	F	6.84	1.46
41	F	3.95	0.84
42	M	17.23	1.32 *
43	M	15.76	1.52 **
44	M	16.64	1.50 ***
19	M	20.81	1.32

**K = Relative Retention time**

**K = Relative Retention time =**

**R<sub>t</sub> amide/R<sub>t</sub> GE 2270 proper  
starting material (i.e. the  
compound of formula II wherein  
W is COOH)**

### EXPERIMENTAL SECTION

#### TABLE IV - N.M.R.

The  $^1\text{H}$ -NMR spectra were recorded with a Bruker spectrometer in  $\text{DMSO-d}_6$  (hexadeuterodimethylsulfoxide) using TMS as the internal standard (0.00 ppm) ( $\delta$ , ppm, m) at 250 MHz and/or 500 MHz (s=singlet, br s = broad singlet, d=doublet, dd=doublet of doublets, t=triplet, m=multiplet)

#### TABLE V - I.R.

The infrared spectra (IR) were recorded with a Perkin Elmer mod. 580 spectrophotometer in nujol mull.

#### TABLE VI - U.V.

The ultraviolet absorption spectra were recorded with a Perkin Elmer Model 320 spectrometer.

It will be clear to the skilled technician that the data represented in TABLES IV, V and VI below, do not represent all the values of the peaks obtained but only the values of those peaks which permit to characterize the single substance.

TABLE IV - N.M.R. Spectra

COMPOUND NO.	<sup>1</sup> H-NMR (DMSO-d <sub>6</sub> ) δ(ppm)
1	0.84 (d, 3H); 0.87 (d, 3H); 2.57 (s, 3H); 3.39 (s, 3H); 3.77 (dd, 1H); 3.99 (d, 2H); 4.25 (dd, 1H); 4.96 (s, 2H); 7.36-7.22 (m, 7H); 8.28 (s, 1H); 8.50 (s, 1H); 8.59 (s, 1H)
2	0.79 (d, 3H); 0.85 (d, 3H); 2.05-1.70 (m, 4H); 2.54 (s, 3H); 3.33 (s, 3H); 3.65 (m, 2H); 3.81 (dd, 1H); 4.10 (m, 1H); 4.35 (dd, 1H); 4.99 (s, 2H); 7.35-7.05 (m, 7H); 8.20 (s, 1H); 8.42 (s, 1H); 8.58 (s, 1H)
3	0.84 (d, 3H); 0.87 (d, 3H); 2.58 (s, 3H); 3.37 (s, 3H); 3.80 (dd, 2H); 3.84 (dd, 1H); 3.91 (dd, 1H); 4.26 (dd, 1H); 4.55 (m, 1H); 4.97 (s, 2H); 7.36-7.20 (m, 7H); 8.29 (s, 1H); 8.55 (s, 1H); 8.59 (s, 1H)
4	0.85 (d, 3H); 0.89 (d, 3H); 2.58 (s, 3H); 2.90 (m, 2H); 3.38 (s, 3H); 3.70 (dd, 1H); 4.29 (dd, 1H); 4.85 (m, 1H); 4.98 (s, 2H); 7.40-7.20 (m, 7H); 8.28 (s, 1H); 8.52 (s, 1H); 8.58 (s, 1H)
5	0.85 (d, 3H); 0.88 (d, 3H); 2.58 (s, 3H); 3.11 (m, 2H); 3.26 (br; s, 1H); 3.38 (s, 3H); 3.78 (dd, 1H); 4.28 (dd, 1H) 4.64 (m, 1H); 4.97 (s, 2H); 6.68 (d, 1H); 7.09 (d, 1H); 7.40-7.20 (m, 7H); 8.27 (s, 1H); 8.47 (s, 1H); 8.59 (s, 1H)

TABLE IV - N.M.R. Spectra (continued)

COMPOUND NO.	<sup>1</sup> H-NMR (DMSO-d <sub>6</sub> ) δ(ppm)
6	0.84 (d, 3H); 0.87 (d, 3H); 0.92 (d, 3H); 0.95 (d, 3H); 1.69 (m, 2H); 1.86 (m, 1H); 2.57 (s, 3H); 3.37 (s, 3H); 3.78 (dd, 1H); 4.26 (dd, 1H); 4.53 (m, 1H); 4.97 (s, 2H); 7.38-7.20 (m, 7H); 8.28 (s, 1H); 8.46 (s, 1H); 8.59 (s, 1H)
7	0.84 (d, 3H); 0.88 (d, 3H); 2.58 (s, 3H); 3.20 (m, 2H); 3.37 (s, 3H); 3.77 (dd, 1H); 4.25 (dd, 1H); 4.73 (m, 1H); 7.40-7.2 (m, 12H); 8.28 (s, 1H); 8.47 (s, 1H) 8.59 (s, 1H)
8	0.85 (d, 3H); 0.89 (d, 3H); 2.08 (s, 3H); 2.16 (m, 2H); 2.56 (m, 2H); 2.57 (s, 3H); 3.40 (s, 3H); 3.79 (dd, 1H); 4.27 (dd, 1H); 4.61 (m, 1H); 5.00 (s, 2H); 7.37-7.20 (m, 7H); 8.29 (s, 1H); 8.52 (s, 1H); 8.60 (s, 1H)
9	0.84 (d, 3H); 0.88 (d, 3H); 2.45-1.70 (m, 4H); 2.58 (s, 3H); 3.37 (s, 3H); 3.68 (m, 2H); 3.78 (dd, 1H); 4.10 (m, 1H); 4.27 (dd, 1H); 4.49 (m, 1H); 7.35-7.22 (m, 7H); 8.27 (s, 1H); 8.50 (s, 1H); 8.59 (s, 1H)
10	0.85 (d, 3H); 0.88 (d, 3H); 1.19 (d, 3H); 2.59 (s, 3H); 3.39 (s, 3H); 3.78 (dd, 1H); 4.30 (m, 2H); 4.48 (dd, 1H); 4.99 (s, 2H); 7.4-7.2 (m, 7H); 8.33 (s, 1H); 8.49 (s, 1H); 8.60 (s, 1H)

TABLE IV - N.M.R. Spectra (continued)

COMPOUND NO.	<sup>1</sup> H-NMR (DMSO-d <sub>6</sub> ) δ(ppm)
11	0.84 (d, 3H); 0.88 (d, 3H); 1.55-1.35 (m, 2H); 1.61 (m, 2H); 1.83 (m, 2H); 2.58 (s, 3H); 3.34 (m, 2H); 3.38 (s, 3H); 3.79 (dd, 1H); 3.91 (br; s, 1H); 4.29 (dd, 1H); 4.97 (s, 2H); 7.35-7.13 (m, 7H); 8.27 (s, 1H); 8.43 (s, 1H); 8.59 (s, 1H)
12	0.85 (d, 3H); 0.88 (d, 3H); 1.37 (m, 2H); 1.70-1.49 (m, 3H); 1.75 (m, 1H); 2.58 (s, 3H); 2.76 (m, 2H); 3.38 (s, 3H); 3.78 (dd, 1H); 4.03 (m, 2H); 4.28 (m, 2H); 4.97 (s, 2H); 7.35-7.20 (m, 7H); 8.28 (s, 1H); 8.49 (s, 1H); 8.59 (s, 1H)
13	0.84 (d, 3H); 0.88 (d, 3H); 1.98-1.82 (m, 2H); 2.18 (m, 2H); 2.56 (s, 3H); 2.69 (dd, 2H); 3.36 (s, 3H); 3.85-3.62 (m, 3H); 4.31 (m, 2H); 4.85 (m, 1H); 4.96 (s, 2H); 7.38-7.19 (m, 7H); 8.24 (s, 1H); 8.55 (s, 1H); 8.63 (s, 1H)
14	0.85 (d, 3H); 0.88 (d, 3H); 1.99-1.82 (m, 3H); 2.06 (m, 1H); 2.58 (s, 3H); 3.58 (m, 1H); 3.67 (m, 1H); 3.79 (dd, 1H); 4.18 (d, 2H); 4.28 (dd, 1H); 4.97 (s, 2H); 6.93 (s, 1H); 7.36-7.28 (m, 8H); 8.28 (s, 1H); 8.52 (s, 1H); 8.59 (s, 1H)
15	0.85 (d, 3H); 0.88 (d, 3H); 2.07-1.63 (m, 4H); 2.58 (s, 3H); 2.99 (dd, 1H); 3.09 (dd, 1H); 3.38 (s, 3H); 3.51 (m, 1H); 3.77 (m, 2H); 4.30 (m, 2H); 4.89 (m, 1H); 4.98 (s, 2H); 6.66 (d, 1H); 6.95 (br; s, 1H); 7.16 (d, 1H); 7.39-7.20 (m, 8H); 8.23 (s, 1H); 8.42 (s, 1H); 8.58 (s, 1H)

TABLE IV - N.M.R. Spectra (continued)

COMPOUND NO.	<sup>1</sup> H-NMR (DMSO-d <sub>6</sub> ) δ(ppm)
16	0.84 (d, 3H); 0.87 (d, 3H); 1.20 (d, 3H); 1.98-1.80 (m, 3H); 2.08 (m, 1H); 2.56 (s, 3H); 3.36 (s, 3H); 3.85-3.71 (m, 2H); 4.13 (m, 1H); 4.28 (dd, 1H); 4.31 (dd, 1H); 4.73 (m, 1H); 5.05 (d, 1H); 6.89 (br; s, 1H); 7.15 (br; s, 1H); 7.38-7.19 (m, 7H); 8.26 (s, 1H); 8.51 (s, 1H); 8.56 (s, 1H)
17	0.84 (d, 3H); 0.87 (d, 3H); 0.94 (d, 3H); 0.98 (d, 3H); 2.10-1.62 (m, 7H); 2.56 (s, 3H); 3.36 (s, 3H); 3.65 (m, 1H); 3.88-3.70 (m, 2H); 4.31 (m, 2H); 4.88 (m, 1H); 4.96 (s, 2H); 6.79 (br, s, 1H); 7.18 (br; s, 1H); 7.35-7.20 (m, 7H); 8.25 (s, 1H); 8.48 (s, 1H); 8.56 (s, 1H)
18	0.84 (d, 3H); 0.88 (d, 3H); 1.81 (m, 2H); 2.30 (t, 2H); 2.58 (s, 3H); 3.35 (m, 2H); 3.37 (s, 3H); 3.78 (dd, 1H); 4.28 (dd, 1H); 4.97 (s, 2H); 7.35-7.20 (m, 7H); 8.27 (s, 1H); 8.46 (s, 1H); 8.59 (s, 1H)
19	0.84 (d, 3H); 0.87 (d, 3H); 1.35 (m, 2H); 1.56 (m, 4H); 2.22 (t, 2H); 2.58 (s, 3H); 3.36 (m, 2H); 3.38 (s, 3H); 3.80 (dd, 1H); 4.29 (dd, 1H); 4.97 (s, 2H); 7.42-7.22 (m, 7H); 8.29 (s, 1H); 8.45 (s, 1H); 8.62 (s, 1H)

TABLE IV - N.M.R. Spectra (continued)

COMPOUND NO.	<sup>1</sup> H-NMR (DMSO-d <sub>6</sub> ) δ(ppm)
20	0.84 (d, 3H); 0.88 (d, 3H); 1.31 (br; s, 6H); 1.51 (m, 2H); 1.57 (m, 2H); 2.19 (t, 2H); 2.58 (s, 3H); 3.32 (m, 2H); 3.37 (s, 3H); 3.79 (dd, 1H); 4.28 (dd, 1H); 4.97 (s, 2H); 7.38-7.19 (m, 7H); 8.27 (s, 1H); 8.45 (s, 1H); 8.59 (s, 1H)
21	0.84 (d, 3H); 0.88 (d, 3H); 1.41-1.20 (m, 12H); 1.47 (m, 2H); 1.57 (m, 2H); 2.17 (t, 2H); 2.58 (s, 3H); 3.29 (m, 2H); 3.38 (s, 3H); 3.79 (dd, 1H); 4.28 (dd, 1H); 4.97 (s, 2H); 7.38-7.19 (m, 7H); 8.27 (s, 1H); 8.43 (s, 1H); 8.59 (s, 1H)
22	0.85 (d, 3H); 0.87 (d, 3H); 2.57 (s, 3H); 2.79 (t, 2H); 3.37 (s, 3H); 3.59 (t, 2H); 3.78 (dd, 1H); 4.28 (dd, 1H); 4.97 (s, 1H); 7.41-7.20 (m, 7H); 8.27 (s, 1H); 8.44 (s, 1H); 8.57 (s, 1H)
23	0.84 (d, 3H); 0.87 (d, 3H); 1.67 (m, 2H); 2.53 (t, 2H); 2.57 (s, 3H); 3.26 (t, 2H); 3.37 (s, 3H); 3.78 (dd, 1H); 4.28 (dd, 1H); 4.97 (s, 2H); 7.41-7.26 (m, 7H); 8.26 (s, 1H); 8.44 (s, 1H); 8.57 (s, 1H)
24	0.85 (d, 3H); 0.88 (d, 3H); 1.58 (m, 2H); 1.79 (m, 2H); 2.58 (s, 3H); 3.38 (s, 3H); 3.50 (m, 2H); 3.78 (dd, 1H); 4.28 (dd, 1H); 4.97 (s, 2H); 7.38-7.21 (m, 7H); 8.27 (s, 1H); 8.45 (s, 1H); 8.59 (s, 1H)



TABLE IV - N.M.R. Spectra (continued)

COMPOUND NO.	<sup>1</sup> H-NMR (DMSO-d <sub>6</sub> ) δ (ppm)
25	0.84 (d, 3H); 0.88 (d, 3H); 1.65-1.35 (m, 8H); 4.58 (s, 3H); 3.38 (s, 3H); 3.78 (dd, 1H); 4.28 (dd, 1H); 4.97 (s, 2H); 7.40-7.20 (m, 7H); 8.28 (s, 1H); 8.43 (s, 1H); 8.59 (s, 1H)
26	0.85 (d, 3H); 0.89 (d, 3H); 2.56 (s, 3H); 3.36 (s, 3H); 3.80 (dd, 1H); 4.31 (dd, 1H); 4.62 (br, s, 2H); 4.96 (s, 2H); 7.39-7.15 (m, 7H); 7.47 (d, 2H); 7.90 (d, 2H); 8.26 (s, 1H); 8.41 (s, 1H); 8.58 (s, 1H)
27	0.84 (d, 3H); 0.88 (d, 3H); 1.62 (br, s, 2H); 1.92 (br, s, 2H); 2.58 (s, 3H); 2.60 (m, 1H); 3.38 (s, 3H); 3.79 (dd, 1H); 4.16 (m, 2H); 4.29 (dd, 1H); 4.38 (m, 2H); 7.35-7.19 (m, 7H); 8.25 (s, 1H); 8.29 (s, 1H); 8.57 (s, 1H)
28	0.85 (d, 3H); 0.89 (d, 3H); 1.39 (m, 2H); 1.61 (m, 2H); 1.76 (m, 2H); 2.58 (s, 3H); 2.88 (t, 2H); 3.33 (m, 2H); 3.80 (dd, 1H); 4.29 (dd, 1H); 4.98 (s, 2H); 7.34-7.20 (m, 7H); 8.26 (s, 1H); 8.45 (s, 1H); 8.58 (s, 1H)
29	0.84 (d, 3H); 0.88 (d, 3H); 2.58 (s, 6H); 3.38 (s, 3H); 3.70-3.41 (m, 5H); 3.89-3.75 (m, 2H); 3.98 (br, s, 1H); 4.35-4.26 (m, 2H); 4.97 (s, 2H); 7.35-7.21 (m, 7H); 8.26 (s, 1H); 8.28 (s, 1H); 8.58 (s, 1H)

TABLE IV - N.M.R. Spectra (continued)

COMPOUND NO.	<sup>1</sup> H-NMR (DMSO-d <sub>6</sub> ) δ(ppm)
30	0.84 (d, 3H); 0.88 (d, 3H); 2.58 (s, 3H); 3.29-3.14 (m, 2H); 3.38 (s, 3H); 3.90-3.49 (m, 4H); 4.29 (dd, 1H); 4.92 (m, 1H); 4.97 (s, 2H); 5.12 (t, 1H); 7.35-7.18 (m, 7H); 8.26 (s, 1H); 8.51 (s, 1H); 8.58 (s, 1H)
31	0.86 (d, 3H); 0.89 (d, 3H); 1.81 (m, 2H); 2.59 (s, 3H); 3.32 (m, 4H); 3.39 (s, 3H); 3.80 (dd, 1H); 4.30 (dd, 1H); 4.99 (s, 2H); 6.75 (d, 1H); 7.41-7.18 (m, 9H); 8.28 (s, 1H); 8.46 (s, 1H); 8.59 (s, 1H)
32	0.85 (d, 3H); 0.88 (d, 3H); 2.21 (s, 6H); 2.59 (s, 3H); 3.38 (s, 3H); 3.43 (m, 4H); 3.81 (dd, 1H); 4.31 (dd, 1H); 4.98 (s, 2H); 7.45-7.19 (m, 7H); 8.28 (s, 1H); 8.45 (s, 1H); 8.61 (s, 1H)
33	0.86 (d, 3H); 0.90 (d, 3H); 1.91-1.70 (m, 2H); 2.26-2.05 (m, 2H); 2.60 (s, 3H); 2.91-2.69 (m, 4H); 3.40 (s, 3H); 3.51 (br; s, 2H); 3.95-3.75 (m, 2H); 4.30 (dd, 1H); 4.99 (s, 2H); 7.41-7.18 (m, 12H); 8.28 (s, 1H); 8.45 (s, 1H); 8.66 (s, 1H)
34	0.85 (d, 3H); 0.89 (d, 3H); 1.81-1.49 (m, 4H); 2.01-1.88 (m, 2H); 2.59 (s, 3H); 2.98-2.65 (m, 4H); 3.39 (s, 3H); 3.80-3.51 (m, 4H); 3.81 (dd, 1H); 4.31 (dd, 1H); 4.99 (s, 2H); 7.41-7.18 (m, 7H); 7.90-7.65 (m, 6H); 8.25 (s, 1H); 8.36 (s, 1H); 8.61 (s, 1H)

TABLE IV - N.M.R. Spectra (continued)

COMPOUND NO.	<sup>1</sup> H-NMR (DMSO-d <sub>6</sub> ) δ (ppm)
35	0.85 (d, 3H); 0.88 (d, 3H); 2.59 (s, 3H); 3.39 (s, 3H); 3.79 (dd, 1H); 4.29 (dd, 1H); 4.98 (s, 2H); 7.40-7.19 (m, 7H); 7.72 (br, s, 1H); 8.03 (br, s, 1H); 8.28 (s, 1H); 8.47 (s, 1H); 8.60 (s, 1H)
36	0.85 (d, 3H); 0.88 (d, 3H); 1.87 (m, 2H); 2.54 (s, 3H); 2.89 (m, 2H); 3.37 (s, 3H); 3.42 (m, 2H); 3.79 (dd, 1H); 4.29 (dd, 1H); 4.98 (s, 2H); 7.38-7.20 (m, 7H); 7.69 (br, s, 3H); 8.29 (s, 1H); 8.49 (s, 1H); 8.61 (s, 1H)
37	0.83 (d, 3H); 0.87 (d, 3H); 1.32 (m, 1H); 2.16 (m, 1H); 2.46 (d, 3H); 2.57 (s, 3H); 2.71 (m, 1H); 3.37 (s, 3H); 3.78 (dd, 1H); 4.16 (d, 1H); 4.26 (dd, 1H); 4.67 (m, 1H); 4.96 (s, 2H); 6.02 (d, 1H); 6.35 (dd, 1H); 7.35-7.20 (m, 7H); 8.28 (s, 1H); 8.49 (s, 1H); 8.60 (s, 1H); 9.61 (s, 1H)
38	0.83 (d, 3H); 0.87 (d, 3H); 1.25 (m, 1H); 2.2 (m, 1H); 2.5 (s, 3H); 2.70 (m, 3H); 3.35 (s, 3H); 3.63 (m, 1H); 3.79 (d, 1H); 4.27 (dd, 1H); 4.97 (s, 2H); 7.4-7.15 (m, 7H); 8.28 (s, 1H); 8.53 (s, 1H); 8.61 (s, 1H)

TABLE IV - N.M.R. Spectra (continued)

COMPOUND NO.	<sup>1</sup> H-NMR (DMSO-d <sub>6</sub> ) δ(ppm)
39	0.83 (d, 3H); 0.87 (d, 3H); 1.32 (m, 1H); 2.16 (m, 1H); 2.47 (d, 3H); 2.57 (s, 3H); 2.72 (m, 4H); 3.37 (s, 3H); 3.50 (m, 2H); 3.78 (dd, 1H); 4.27 (dd, 1H); 4.97 (s, 2H); 7.40-7.20 (m, 7H); 8.28 (s, 1H); 8.49 (s, 1H); 8.60 (s, 1H)
40	0.83 (d, 3H); 0.87 (d, 3H); 1.32 (m, 1H); 1.71 (m, 2H); 2.25-2.14 (m, 3H); 2.46 (d, 3H); 2.57 (s, 3H); 2.7 (m, 1H); 3.37 (s, 3H); 3.76 (dd, 1H); 4.27 (dd, 1H); 4.97 (s, 2H); 5.81 (d, 1H, J=15.7 Hz); 6.78 (m, 1H); 7.39-7.12 (m, 7H); 8.28 (s, 1H); 8.45 (s, 1H); 8.60 (s, 1H)
41	0.86 (d, 3H); 0.89 (d, 3H); 1.43 (m, 1H); 2.19 (m, 1H); 2.47 (d, 3H); 2.59 (s, 3H); 2.72 (m, 1H); 3.39 (s, 3H); 3.50 (t, 2H); 3.58 (t, 2H); 3.68 (t, 2H); 3.79 (dd, 1H); 4.99 (s, 2H); 7.42-7.20 (m, 7H); 8.27 (s, 1H); 8.47 (s, 1H); 8.59 (s, 1H)
42	0.83 (d, 3H); 0.85 (d, 3H); 1.2-1.4 (m, 3H); 1.5-1.65 (m, 4H); 2.22 (t, 3H); 2.60 (d, 1H); 2.69 (d, 1H); 3.37 (s, 3H); 3.79 (dd, 1H); 4.27 (dd, 1H); 4.86 (d, 2H); 4.97 (s, 2H); 5.00 (dd, 1H); 5.1-5.4 (m, 3H); 5.74 (t, 1H); 6.00 (d, 1H); 7.2-7.4 (m, 7H); 8.27 (s, 1H); 8.44 (s, 1H); 8.62 (s, 1H)

TABLE IV - N.M.R. Spectra (continued)

COMPOUND NO.	<sup>1</sup> H-NMR (DMSO-d <sub>6</sub> ) δ(ppm)
43	0.84 (d, 3H); 0.89 (d, 3H); 1.4-1.2 (m, 3H); 1.65-1.50 (m, 4H); 2.23 (t, 3H); 2.59 (s, 3H); 2.79 (m, 1H); 3.87 (m, 1H); 4.25 (m, 1H); 5.04 (t, 1H); 5.35-5.20 (m, 3H); 6.09 (d, 1H); 6.67 (br, s, 1H); 7.04 (br, s, 1H); 7.35-7.15 (m, 6H); 8.24 (s, 1H); 8.26 (s, 1H); 8.45 (s, 1H); 8.61 (s, 1H)
44	0.84 (d, 3H); 0.88 (d, 3H); 1.4-1.25 (m, 3H); 1.65-1.50 (m, 4H); 2.23 (t, 3H); 2.58 (s, 3H); 2.75 (m, 1H); 3.78 (dd, 1H); 4.28 (dd, 1H); 4.98 (m, 3H); 5.35-5.15 (m, 3H); 6.03 (m, 2H); 7.42-7.15 (m, 7H); 8.30 (s, 1H); 8.45 (s, 1H); 8.62 (s, 1H)

TABLE V - I.R. Spectra

COMPOUND NO.	I.R. (nujol cm <sup>-1</sup> )
1	3370; 3110; 1730; 1655; 1545; 1520
2	3350; 3110; 1720; 1650; 1535; 1500
3	3340; 3105; 1720; 1645; 1535; 1500
4	3360; 1725; 1640; 1535
5	3350; 3110; 1725; 1650; 1535; 1510
6	3370; 3105; 1725; 1655; 1535; 1500
7	3360; 3100; 1725; 1655; 1535; 1490
8	3370; 3105; 1725; 1655; 1535; 1505
9	3370; 3110; 3100; 1725; 1657; 1550; 1530; 1505
10	3370; 3105; 1730; 1655; 1540; 1510
11	3359; 3115; 1653; 1551; 1510;
12	3360; 3113; 1720; 1662; 1547; 1510

TABLE V - I.R. Spectra (continued)

COMPOUND NO.	I.R. (nujol cm <sup>-1</sup> )
13	3370; 3110; 1720; 1655; 1530; 1505
14	3350; 3120; 1655; 1535; 1500
15	3350; 3100; 1650; 1530; 1510;
16	3340; 3105; 1650; 1530
17	3340; 3100; 1655; 1530
18	3350; 3100; 1710; 1645; 1540
19	3360; 3115; 1720; 1665; 1540; 1506;
20	3350; 3113; 1720; 1659; 1549; 1506
21	3340; 1710; 1645; 1540; 1500
22	3304; 1653; 1540
23	3333; 1657; 1547; 1092; 1038;
24	3354; 3113; 1653; 1550; 1506; 1245

TABLE V - I.R. Spectra (continued)

COMPOUND NO.	I.R. (nujol cm <sup>-1</sup> )
25	3348; 3111; 1660; 1548; 1507; 1245
26	3315; 1653; 1539; 1238
27	3361; 3113; 1720; 1653; 1531; 1507; 1092
28	3333; 1653; 1547; 1494; 1243
29	3356; 3114; 1653; 1508; 1088
30	3360; 1670; 1505; 1200
31	3351; 3115; 1653; 1549; 1509; 1250
32	3370; 3110; 1655; 1545; 1500; 1245
33	3350; 1655; 1530; 1490; 1220
34	3360; 3105; 1650; 1545; 1510; 1240
35	3320; 1747; 1650; 1540; 1225
36	3330; 1662; 1547; 1496; 1201



TABLE V - I.R. Spectra (continued)

COMPOUND NO.	I.R. (nujol cm <sup>-1</sup> )
37	3327; 1730; 1653; 1464; 1377
38	3355; 1720; 1657; 1543; 1377
39	3321; 1717; 1652; 1545
40	3337; 1665; 1549
41	3341; 1721; 1653; 1548; 1377
42	3335; 1722; 1647; 1543
43	3317; 1665; 1539
44	3317; 1720; 1649; 1545

TABLE VI  
U.V. DATA  $\lambda_{\max}$  ( $E^{1\%}_{1\text{cm}}$ )

Compound No.	MeOH	HCl 0.1N	Phosphate Buffer pH 7.38	KOH 0.1N
1	309 (290.9)	312	309 (247.9)	309 (252.7)
2	309 (257.5)	310 (222.6)	311	309 (226.3)
3	309 (297.5)	312	309 (229.6)	309 (235.8)
4	309 (245.1)	312	308 (234.7)	308 (234.1)
5	308 (173.8)	312	309 (150.0)	305 (181.2)
6	309 (277.1)	313	309 (229.8)	309 (236.7)
7	309 (258.6)	313	309 (207.9)	309 (218.9)
8	309 (279.8)	311	309 (225.9)	309 (229.4)
9	309 (261.9)	313	308 (228.1)	309 (235.0)
10	309 (279.3)	314	309 (241.8)	309 (251.1)
11	309 (216.8)	310 (178.5)	312	310 (194.9)
12	309 (226.2)	310 (188.3)	311	309 (202.4)

TABLE VI (continued)  
U.V. DATA  $\lambda_{\text{max}}$  ( $E^{1\%}_{1\text{cm}}$ )

Compound No.	MeOH	HCl 0.1N	Phosphate Buffer pH 7.38	KOH 0.1N
13	308 (237.9)	314	308 (247.4)	308 (260.3)
14	309 (263.4)	313	313	314
15	309 (222.6)	313	314	304 (169.8)
16	309 (235.6)	313	312	312
17	309 (230.3)	312	312	312
18	309 (288.8)	313	309 (239.4)	309 (248.0)
19	309 (283.2)	312	309 (220.3)	309 (230.1)
20	309	314	309	309
21	309 (271.6)	313	311 (221.6)	309 (221.6)
22	309 (190.9)	309 (152.2)	308 (160.4)	309 (165.6)
23	309 (242.2)	310 (182.2)	309 (200.9)	309 (200.9)
24	309	312	310	309

TABLE VI (continued)  
U.V. DATA  $\lambda_{\text{max}}$  ( $E_{1\%}^{1\text{cm}}$ )

Compound No.	MeOH	HCl 0.1N	Phosphate Buffer pH 7.38	KOH 0.1N
25	309	312	310	309
26	309 (260.0)	313	310 (197.7)	310 (208.6)
27	310 (264.6)	313	310 (227.4)	310 (232.1)
28	309 (260.5)	314	310 (186.8)	310 (203.6)
29	309 (243.4)	311	312	311
30	309 (248.5)	311	311	309
31	305 (253.7)	310	310	313 (249.5)
32	309 (267.9)	310 (234.9)	312	312
33	309 (247.5)	311 (234.4)	314	312
34	310 (224.0)	309 (198.2)	310	312
35	308 (269.8)	314	313	313
36	309 (243.9)	309 (205.5)	312	313

TABLE VI (continued)  
U.V. DATA  $\lambda_{\text{max}}$  (E<sup>1%</sup><sub>1cm</sub>)

Compound No.	MeOH	HCl 0.1N	Phosphate Buffer pH 7.38	KOH 0.1N
37	309 (255.1)	312	314	312
38	308	308	308	308
39	308 (247.9)	312	308 (201.3)	308 (215)
40	309 (304.3)	312	309 (235.9)	309 (262.0)
41	309 (256.4)	312	309 (215.1)	309 (228.6)
42	309	312	309	307
43	309 (253.6)	313	309 (208.4)	309 (235.4)
44	309 (264.9)	314	309 (208.1)	309 (223.7)

The antimicrobial activity of the compounds of the invention can be demonstrated by a series of standard tests in vitro.

5

MIC for Propionibacterium acnes, and Bacteroides fragilis are determined by agar dilution (inocula  $10^4/10^5$  CFU/spot). MIC for other organisms are determined by microbroth dilution (inocula  $10^4$  to  $10^5$  CFU/ml). Incubation times are 18-24 h, except for Haemophilus influenzae, P. acnes, B. fragilis (48 h). All organisms are incubated at 37°C; H. influenzae is incubated in a 5% CO<sub>2</sub> atmosphere, anaerobes in an anaerobic gas mixture. Media used are: Iso-Sensitest broth (Oxoid) (Staphylococci, Streptococcus faecalis, Escherichia coli, Proteus vulgaris; brain heart infusion broth (Difco) + 1% Supplement C (Difco) (H. influenzae);

20

The minimal inhibitory concentrations (MIC, microgram/ml) for some microorganisms are reported below in Table VII.

25

30



TABLE VII - (MIC, microgram/ml) (continued)

STRAIN	COMPOUND OF EXAMPLE No.					
	14	18	19	20	26	
<u>Staph. aureus</u> L165 Tour	0.06	0.25	0.06	0.25	0.25	
<u>Staph. epidermidis</u> L147 ATCC 12228	0.13	0.25	0.06	0.25	0.25	
<u>Staph. haemolyticus</u> L602	0.25	1	0.25	0.25	0.5	
<u>Strep. pneumoniae</u> L44 UC41	>128	1	0.25	2	2	
<u>Strep. faecalis</u> L149 ATCC 7080	0.06	0.13	0.06	<0.13	0.13	
<u>Prop. acnes</u> L1014 ATCC 6919	0.03	0.016	0.008	0.03	0.008	
<u>Bact. fragilis</u> L1010 ATCC 23745	>128	2	4	>128	32	
<u>Haemophilus Influenzae</u> type B	>128	2	2	>128	8	
<u>Esch. coli</u> L47 SKF 12140	>128	>128	>128	>128	>128	
<u>Prot. vulgaris</u> ATCC 881	>128	>128	>128	>128	>128	



TABLE VII - (MIC, microgram/ml) (continued)

STRAIN	COMPOUND OF EXAMPLE No.			
	27	28	32	35
<u>Staph. aureus</u> L165 Tour	0.25	0.13	0.5	0.13
<u>Staph. epidermidis</u> L147 ATCC 12228	0.5	0.5	0.5	0.13
<u>Staph. haemolyticus</u> L602	1	1	0.5	0.5
<u>Strep. pneumoniae</u> L44 UC41	8	2	1	>128
<u>Strep. faecalis</u> L149 ATCC 7080	1	0.06	0.25	0.06
<u>Prop. acnes</u> L1014 ATCC 6919	0.03	0.008	0.13	0.004
<u>Bact. fragilis</u> L1010 ATCC 23745	64	>128	>128	>128
<u>Haemophilus influenzae</u> type B	8	>128	>128	>128
<u>Esch. coli</u> L47 SKF 12140	>128	>128	>128	>128
<u>Prot. vulgaris</u> ATCC 881	>128	>128	>128	>128

TABLE VII - (MIC, microgram/ml) (continued)

STRAIN	COMPOUND OF EXAMPLE No.				
	36	37	38	39	40
<u>Staph. aureus</u> L165 Tour	0.13	0.03	32	0.13	0.25
<u>Staph. epidermidis</u> L147 ATCC 12228	0.13	0.06	32	0.25	0.25
<u>Staph. haemolyticus</u> L602	0.13	0.13	64	2	1
<u>Strep. pneumoniae</u> L44 UC41	4	>128	>128	4	1
<u>Strep. faecalis</u> L149 ATCC 7080	0.13	0.06	16	≤0.13	≤0.13
<u>Prop. acnes</u> L1014 ATCC 6919	0.06	0.004	0.25	≤0.13	-
<u>Bact. fragilis</u> L1010 ATCC 23745	>128	>128	>128	8	4
<u>Haemophilus Influenzae</u> type B	>128	>128	>128	1	1
<u>Esch. coli</u> L47 SKF 12140	>128	>128	>128	>128	>128
<u>Prot. vulgaris</u> ATCC 881	>128	>128	>128	>128	>128

TABLE VII - (MIC, microgram/ml) (continued)

STRAIN	COMPOUND OF EXAMPLE No.			
	41	42	43	44
<u>Staph. aureus</u> L165 Tour	0.25	0.13	0.13	0.13
<u>Staph. epidermidis</u> L147 ATCC 12228	0.5	0.13	0.5	0.5
<u>Staph. haemolyticus</u> L602	1	0.5	0.5	1
<u>Strep. pneumoniae</u> L44 UC41	2	1	0.5	1
<u>Strep. faecalis</u> L149 ATCC 7080	≤0.13	0.13	0.13	0.06
<u>Prop. acnes</u> L1014 ATCC 6919	≤0.13	0.016	0.016	0.016
<u>Bact. fragilis</u> L1010 ATCC 23745	4	>128	>128	>128
<u>Haemophilus Influenzae</u> type B	1	4	>128	>128
<u>Esch. coli</u> L47 SKF 12140	>128	>128	>128	>128
<u>Prot. vulgaris</u> ATCC 881	>128	>128	>128	>128

In view of their properties, the compounds of the invention can be used as active ingredients in the preparation of medicaments for human or animal treatment

5

In particular, the amide derivatives of the antibiotic GE 2270 compounds of formula I are antimicrobial agents mainly active against gram positive bacteria and gram positive as well as gram negative anaerobes.

10

The main therapeutic indication of the antibiotic substances of the invention is thus in the treatment of infections related to the presence of microorganisms susceptible to them.

15

The term "treatment" is intended to encompass also prophylaxis, therapy and cure.

The patient receiving this treatment is any animal in need, including primates, in particular humans, and other mammals such as equines, cattle, swine and sheep; and poultry and pets in general.

20

The compounds of the invention can be administered as such or in admixture with pharmaceutically acceptable carriers and can also be administered in conjunction with other antimicrobial agents. Conjunctive therapy, thus includes sequential, simultaneous and separate administration of the active compounds in a way that the therapeutical effects of the first administered one is not entirely disappeared when the subsequent is administered.

25

30

A preferred pharmaceutical formulation is represented by a formulation suitable for a topical application on an intact or damaged skin or mucous membrane. Examples of such formulations are powders, ointments, creams and lotions. The excipients in these

formulations are the usual pharmaceutically acceptable vehicles such oleaginous ointment bases (e.g. cetyl esters wax, oleic acid, olive oil, paraffin, spermaceti, starch glycerite); absorbent ointment bases (e.g. anhydrous lanolin, hydrophilic petrolatum), emulsion ointment bases (e.g. cetyl alcohol, glyceryl monostearate, lanolin, stearic acid), water-soluble ointment bases (e.g. glycol ethers and their derivatives which include polyethylene glycols, poly(oxy-1,2-ethanediyl)-alpha-hydro-omega-hydroxy-octadecanoate, polysorbates, and polyethylene glycols mono-stearates).

These formulations may contain other known excipients, such as preservatives and are prepared as known in the art and reported in reference handbooks such as Remington's Pharmaceutical Sciences, Seventeenth edition, 1985, Mack Publishing Co.

The compounds of the invention can also be formulated into formulation suitable for parenteral administration according to procedures known per se in the art. For instance, a compound of the invention is formulated with polypropylene glycol or dimethylacetamide and a surface-active agent such as polyoxyethylene sorbitan mono-oleate or polyethoxylated castor oil.

A preferred formulation for parenteral administration includes the following excipients: Cremophor® EL (polyoxyl 35 castor oil USP/NF) 20%, propylene glycol 5-10%.

Preferably, this formulation is used for i.v. administration in the treatment of any infection involving a microorganism susceptible to an antibiotic of the invention.

An example of a suitable formulation used for I.V. is the following

	compound No. 19	100 mg
5	propylene glycol	1 ml
	water for injection q.s.	5 ml
	phosphate buffer pH 8-8.5	

10

In the treatment of pseudomembranous colitis or other diseases attributable to the presence of anaerobes in the gastrointestinal tract, an effective dose of the compounds of the invention may be administered orally in suitable pharmaceutical form such as a capsule or an aqueous suspension.

The dosage of the active ingredient depends on many factors which include type, age and conditions of the patient, specific active ingredient and formulation selected for administration, administration schedule, etc.

In general, effective antimicrobial dosages are employed per single unit dosage form. Repeated applications of these dosage forms, e.g. from 2 to 6 times a day, are in general preferred. An effective dosage may be in general in the range 0.5-50 mg/kg body weight/day.

A preferred topic preparation is an ointment containing from 1% to 10% of a compound of the present invention.

Anyway, the prescribing physician will be able to determine the optimal dosage for a given patient in a given situation.

5 Besides their use as medicaments in human and veterinary therapy, the compounds of the invention can also be used as animal growth promoters.

10 For this purpose, a compound of the invention is administered orally in a suitable feed. The exact concentration employed is that which is required to provide for the active agent in a growth promotant effective amount when normal amounts of feed are consumed.

15 The addition of the active compound of the invention to animal feed is preferably accomplished by preparing an appropriate feed premix containing the active compound in an effective amount and incorporating the premix into the complete ration.

20 Alternatively, an intermediate concentrate or feed supplement containing the active ingredient can be blended into the feed. The way in which such feed premixes and complete rations can be prepared and administered are described in reference books (such as "Applied Animal Nutrition", W.H. Freedman and CO., S. 25 Francisco, USA, 1969 or "Livestock Feeds and Feeding" O and B books, Corvallis, Oregon, USA, 1977).

30 The following examples further illustrate the invention and should not be interpreted as limiting it in any way.

### EXAMPLES OF THE INVENTION

5            PROCEDURE A - Reaction of GE 2270 factor A<sub>3</sub>  
starting material with the selected amine

10           Example 1:  
Preparation of compound no. 15, 29, 30, 32, 33

To a stirred solution of 1 mmol of GE 2270 factor A<sub>3</sub> (prepared as described in European Patent Application Publication No. 406745) in 10 ml of  
15 dimethylformamide (DMF), 1.2 mmols of the selected amine, 1.4 mmols of triethylamine (TEA) and 1.2 mmols of di-phenylphosphorazidate (DPPA) were added at 0°C. (If the salt (chloride, p-toluenesulfonate, etc) of the  
20 selected amine was used, a double amount of TEA had to be used). The temperature was allowed to rise to room temperature and stirring was continued for about 4 h. The reaction mixture was then acidified with 1N aq HCl to about pH 3 and then diluted with water to complete  
25 precipitation of the product. The wet solid was dried in air and then purified by flash chromatography on silica gel 60 (230 - 400 mesh ASTM - Merck) eluting with 3 to 5% methanol in chloroform. Fractions containing the  
30 title compound were pooled together and the solvent evaporated. Trituration of the solid with ethyl ether yielded the title compound as a fine powder.

PROCEDURE A1 - Reaction of GE 2270 factor A<sub>3</sub>  
starting material with the selected amine containing



further reactive functional group(s), all of which protected, and subsequent deprotection of the protecting group(s).

5

Example 2:

Preparation of compound no. 34, 36

10           The reaction was substantially carried out as described in Example 1. Once the reaction product had been purified by flash chromatography, 1 mmol of the solid obtained was treated with 7 ml of cold  
15           trifluoroacetic acid (TFA). The suspension was swirled for a few minutes until a solution was obtained and TFA was evaporated "in vacuo" in the cold. The gummy product still containing traces of TFA was then treated with  
20           ethyl ether and the title compound was obtained as the trifluoroacetate salt in the form of a fine powder.

Example 3:

Preparation of compound no. 1, 3 to 10, 18 to  
25           21, 39

30           The reaction was substantially carried out as described in Example 1. Once the reaction product had been purified by flash chromatography, 1 mmol of the solid obtained was dissolved in 20 ml of dioxane and 1.2 ml of 1N aq NaOH were added under stirring at room temperature. After 5 h the solution was acidified with 1N aq HCl to pH 2 and diluted with water to complete precipitation of the title compound which was filtered off and allowed to dry in air yielding the title compound as a fine powder.

Example 4:

5

## Preparation of compound no. 2

The reaction was carried out as described in Example 3. Once hydrolysis of the ester function had been accomplished and the compound had been allowed to dry in air, 1 mmols of the solid obtained was dissolved in 20 ml of TFA and 50 mmols of thioanisole were added under stirring at room temperature as described by Y. Kiso et al., Chem. Pharm. Bull. 28, 673, 1980. After 3.5 h, TFA was evaporated "in vacuo" in the cold and the residue taken up in a minimum amount of 1% methanol in chloroform. Addition of ethyl ether induced the precipitation of the title compound which was filtered, washed with more ethyl ether and dried "in vacuo" to yield the trifluoroacetate salt of the title compound as a fine powder.

Example 4bis:

25

## Preparation of compound no. 37

The reaction was substantially carried out as described in Example 1. Once the starting material had disappeared from the reaction mixture, water was added and the precipitate obtained was filtered off, washed with additional water and allowed to dry in air. The crude material was then dissolved in 3 ml of THF and stirred overnight at room temperature in the presence of 10% aq. HCl. Dilution with water provided complete precipitation of the product which was filtered off and allowed to dry in air. The solid was then purified by

flash chromatography on silica gel 60 (230 - 400 mesh ASTM - Merck) eluting with 2 to 4% methanol in chloroform. Fractions containing the title compound were pooled together and the solvent evaporated yielding a pale yellow powder.

**PROCEDURE B - Reaction of GE 2270 factor A<sub>3</sub>**  
starting material with the selected amine containing unprotected acid moieties.

**Example 5**  
Preparation of compound no. 19, 22 to 28, 40, 41

1.1 mmols of DPPA were added at 0°C to a stirred solution of 1 mmol of GE 2270 factor A<sub>3</sub> and 1.5 mmols of TEA in 10 ml of DMF. The temperature was allowed to rise to room temperature and stirring was continued for 4.5 more hours. 1.5 mmols of the selected amine and 2 mmols of TEA were then added to the solution at room temperature and stirring was continued at the same temperature for 5 more hours. (If the selected amine contained more than one acid function, the amount of TEA was adjusted so to free the amino group). The reaction mixture was then acidified with 1N aq HCl to about pH 2 and then diluted with water to complete precipitation of the product. The wet solid was dried in air and then purified by flash chromatography on silica gel 60 (230 - 400 mesh ASTM - Merck) eluting with 5 to 10% methanol in chloroform. Fractions containing the title compound were pooled together and the solvent evaporated. Trituration

of the solid with ethyl ether yielded the title compound as a fine powder.

5

PROCEDURE B1 - Reaction of GE 2270 factor A<sub>3</sub> starting material with the selected amine containing reactive functional group(s), all of which are variously protected, in addition to the unprotected acid group(s) and subsequent deprotection of the protecting group(s).

15

Example 6:

Preparation of compound no. 11, 12

The reaction was substantially carried out as described in Example 5. Once the reaction product had been purified by flash chromatography, 1 mmol of the solid obtained was dissolved in 20 ml of TFA and 50 mmols of thioanisole were added under stirring at room temperature. After 3.5 h, TFA was evaporated "in vacuo" in the cold and the residue taken up in a minimum amount of 1% methanol in chloroform. Addition of ethyl ether induced the precipitation of the title compound which was filtered, washed with more ethyl ether and dried "in vacuo" to yield the trifluoroacetate salt of the title compound as a fine powder.

PROCEDURE C - Reaction of selected amide derivatives of GE 2270 factor A<sub>3</sub> as starting material with the selected reagent.

Example 7

5 Preparation of compound no. 14, 15, 16, 17 from  
compound no. 1, 5, 10, 6 respectively

To a stirred solution of 1 mmol of the  
appropriate amide derivative of GE 2270 factor A<sub>3</sub>  
10 (prepared as described in the previous examples) in 10  
ml of DMP, 1.2 mmols of the selected amine, 1.4 mmols of  
TEA and 1.2 mmols of DPPA were added at 0°C. (If the  
salt (chloride, p-toluenesulfonate, etc) of the selected  
15 amine was used, a double amount of TEA had to be used).  
The temperature was allowed to rise to room temperature  
and stirring was continued for about 4 h. The reaction  
mixture was then acidified with 1N aq HCl to about pH 3  
and then diluted with water to complete precipitation of  
20 the product. The wet solid was dried in air and then  
purified by flash chromatography on silica gel 60 (230 -  
400 mesh ASTM - Merck) eluting with 3 to 5% methanol in  
chloroform. Fractions containing the title compound were  
pooled together and the solvent evaporated. Trituration  
25 of the solid with ethyl ether yielded the title compound  
as a fine powder.

30 PROCEDURE C1 - Reaction of the selected amide  
derivative of the GE 2270 factor A<sub>3</sub> as starting material  
with the selected reagent which contains further  
reactive functional group(s), all of which protected,  
and subsequent deprotection of the protecting group(s).

Example 8:

Preparation of compound no. 13 from compound  
no.3

5

The reaction was carried out as described in  
Example 7. Once the reaction product had been purified  
by flash chromatography, 1 mmol of the solid obtained  
was dissolved in 20 ml of dioxane and 1.2 ml of 1N aq  
10 NaOH were added under stirring at room temperature.  
After 5 h the solution was acidified with 1N aq HCl to  
pH 2 and diluted with water to complete precipitation of  
the title compound which was filtered off and allowed to  
dry in air yielding the title compound as a fine powder.  
15

Example 9:

Preparation of compound no. 31 from compound  
no. 36  
20

To a stirred solution of 1 mmol of the  
appropriate amide derivative of GE 2270 factor A<sub>3</sub>  
(prepared as described in the previous examples) in 10  
25 ml of 10% methanolic chloroform, 1.2 mmols of TEA and  
1.1 mmols the selected reagent (see table -) were added  
at room temperature. After 20 min the solvent was  
evaporated "in vacuo" and the residue treated with 5% aq  
Na<sub>2</sub>CO<sub>3</sub>. The solid obtained was filtered off, washed with  
30 more 5% Na<sub>2</sub>CO<sub>3</sub> and water and finally redissolved in 10  
ml of methanol. To this solution, 0.5 ml of water and  
0.1 mmols of p-toluenesulfonic acid were added and the  
reaction mixture was stirred at room temperature  
overnight. The solution was then reduced to a small  
volume (about 2 ml) under vacuum and water was added to

precipitate the title compound which, after drying in air, was obtained as a fine powder.

5

Example 9bis:

Preparation of compound no. 38 from compound no. 37

10

To a stirred solution of 0.23 mmols of the appropriate amide derivative of GE 2270 factor A<sub>3</sub> (prepared as described in the previous examples) in 40 ml of ethanol, 9.2 mmols of acetic acid, 9.2 mmols of sodium acetate and 0.506 mmols of the selected reagent (see table II) were added at room temperature. After 15 2 hours 0.46 mmols of NaBH<sub>4</sub> (Fluka) were added and stirring was continued overnight at the same temperature. Evaporation of the solvent provided a crude 20 material which was washed with 10 ml of 1N HCl, filtered and allowed to dry in air. The solid was then purified by flash chromatography on silica gel 60 (230 - 400 mesh ASTM - Merck) eluting with 0 to 10% methanol in dichloromethane. The fractions containing the methyl 25 ester of the title compound (intermediate) were pooled together and the solvent evaporated providing a solid which was redissolved in 2 ml dioxane and treated overnight with a 1.2 molar excess of 1N NaOH at room temperature. Evaporation of the solvent gave a solid 30 which was further purified by trituration with a 1:1 mixture of ethyl acetate:methanol yielding the title compound as a fine powder.

PROCEDURE D - Reaction of GE 2270 factor A<sub>2</sub>  
starting material with the selected amine

5

Example 10:

Preparation of compound No. 35

1 mmol of GE 2270 factor A<sub>2</sub> (prepared as  
described in European Patent Application Publication  
No. 406745) were dissolved in 10 ml of a saturated solu-  
tion of methanolic ammonia. The solution was allowed to  
stand for 3 days at room temperature and then evaporated  
"in vacuo". The residue was taken up in 2 ml of methanol  
and the title compound precipitated with water, filtered  
off and allowed to dry in air. Trituration with ethyl  
ether yielded the title compound as a fine powder.

20

PROCEDURE E - Preparation of a salt of a  
compound of the invention.

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Example 11:

Preparation of the arginine salt of  
compound No. 19

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To a suspension of 3 g of compound No. 19  
(2.42 mmols) 180 ml of dioxane, a solution of 423 mg of  
L-arginine (2.42 mmols) in 120 ml of water were added  
under stirring and the non clear solution was thus  
lyophilized to recovered the desired salt.



5                    PROCEDURE F - Reaction of GE 2270 component C<sub>2a</sub>  
starting material (i.e. the compound of formula II  
wherein R is methoxymethyl, R<sub>1</sub> is methyl, R<sub>4</sub> is  
hydroxymethyl and W is COOH) with the selected amine  
containing further reactive functional group(s), all of  
which protected, and subsequent deprotection of the  
protecting group(s).

10

Example 12:

Preparation of compound no. 42

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The reaction was carried out as described in  
Example 3 using GE 2270 component C<sub>2a</sub> starting material  
instead of factor A<sub>3</sub>.

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PROCEDURE G - Reaction of GE 2270 component C<sub>2a</sub>  
starting material as described in procedure F with the  
selected amine containing unprotected acid moieties.

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Example 13:

Preparation of compound no.42

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The reaction was carried out as described in  
Example 5 using GE 2270 component C<sub>2a</sub> starting material  
instead of factor A<sub>3</sub>.

PROCEDURE H - Reaction of GE 2270 component D<sub>1</sub> starting material (i.e. the compound of formula II wherein R and R<sub>1</sub> are hydrogen, R<sub>4</sub> is methyl and W is COOH) with the selected amine containing further reactive functional group(s), all of which protected, and subsequent deprotection of the protecting group(s).

Example 14:  
Preparation of compound no. 43

The reaction was carried out as described in Example 3 using GE 2270 component D<sub>1</sub> starting material instead of factor A<sub>3</sub>.

PROCEDURE I - Reaction of GE 2270 component D<sub>1</sub> starting material as described in procedure H with the selected amine containing unprotected acid moieties.

Example 15:  
Preparation of compound no.43

The reaction was carried out as described in Example 5 using GE 2270 component D<sub>1</sub> starting material instead of factor A<sub>3</sub>.

PROCEDURE J - Reaction of GE 2270 component D<sub>2</sub> (i.e. the compound of formula II wherein R is hydroxymethyl, R<sub>1</sub> and R<sub>4</sub> are methyl and W is COOH)

starting material with the selected amine containing further reactive functional group(s), all of which protected, and subsequent deprotection of the protecting group(s).

Example 16:

Preparation of compound no. 44

The reaction was carried out as described in Example 3 using GE 2270 component D<sub>2</sub> starting material instead of factor A<sub>3</sub>.

PROCEDURE K - Reaction of GE 2270 component D<sub>2</sub> starting material as described in procedure J with the selected amine containing unprotected acid moieties.

Example 17:

Preparation of compound no.44

The reaction was carried out as described in Example 5 using GE 2270 component D<sub>2</sub> starting material instead of factor A<sub>3</sub>.

PROCEDURE L - Reaction of a mixture of minor components (C<sub>2a</sub>, D<sub>1</sub>, D<sub>2</sub> and E) of antibiotic GE 2270 (starting material) with the selected amine containing further reactive functional group(s), all of which

protected, and subsequent deprotection of the protecting group(s).

5

Example 18:

The reaction was carried out as described in Example 3 using a mixture of minor components (C<sub>2a</sub>, D<sub>1</sub>, D<sub>2</sub> and E) of antibiotic GE 2270 starting material instead of factor A<sub>3</sub> and methyl 6-aminocaproate hydrochloride (Fluka). R<sub>t</sub> (min) refer to HPLC method M reported in the HPLC analysis section.

When Y=--NH CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>COOCH<sub>3</sub>, R<sub>t</sub> (min) are respectively 43.43 for GE 2270 factor C<sub>2a</sub>, 39.42 for GE 2270 factor D<sub>1</sub>, 42.29 for GE 2270 factor D<sub>2</sub> and 37.41 for GE 2270 factor E.

When Y=--NH CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>COOH, R<sub>t</sub> (min) are respectively 17.23 for GE 2270 factor C<sub>2a</sub>, 15.76 for GE 2270 factor D<sub>1</sub>, 16.64 for GE 2270 factor D<sub>2</sub> and 15.13 for GE 2270 factor E.

25

PROCEDURE M - Reaction of a selected mixture of minor components (C<sub>2a</sub>, D<sub>1</sub>, D<sub>2</sub> and E) of antibiotic GE 2270 starting material with the selected amine containing unprotected acid moieties.

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Example 19:

The reaction was carried out as described in Example 5 using a selected mixture of minor components (C<sub>2a</sub>, D<sub>1</sub>, D<sub>2</sub> and E) of antibiotic GE 2270 starting

material instead of factor A<sub>3</sub> and 6-aminocaproic acid (Fluka).

5 R<sub>t</sub> (min) refer to Method M reported in the HPLC analysis section and are respectively 17.23 for GE 2270 factor C<sub>2a</sub>, 15.76 for GE 2270 factor D<sub>1</sub>, 16.64 for GE 2270 factor D<sub>2</sub> and 15.13 for GE 2270 factor E.

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PREPARATION OF THE STARTING MATERIALS

- 5  
1. The following starting materials have been  
purchased from Fluka (Fluka,  
Chemika-Biochemika, Buchs, Switzerland):  
Glycine ethyl ester hydrochloride,  
10 L-threonine methyl ester hydrochloride,  
L-tyrosine methyl ester hydrochloride,  
L-leucine methyl ester hydrochloride,  
L-phenylalanine methyl ester hydrochloride,  
L-methionine methyl ester hydrochloride,  
15 L-proline methyl ester hydrochloride,  
L-threonine methyl ester hydrochloride,  
Na-Cbz-L-lysine,  
methyl 4-aminobutyrate hydrochloride,  
20 methyl 6-aminocaproate hydrochloride,  
6-aminocaproic acid,  
4-(methylamino)benzoic acid,  
piperidine-4-carboxylic acid,  
N-methyl-D-glucamine,  
25 D(+)-glucosamine hydrochloride,  
2-dimethylaminoethylamine,  
amino acetaldehyde dimethylacetal,  
 $\beta$ -alanine ethyl ester hydrochloride.
- 30  
2. The following starting materials have been  
purchased from Sigma (Sigma, Biochemicals  
Organic Compounds, St. Louis, U.S.A.):  
N $\epsilon$ -Cbz-L-ornithine,  
L-aspartic acid dimethyl ester hydrochloride.

3. The following starting materials have been purchased from Aldrich (Aldrich, Catalogo Prodotti di Chimica Fine, Milano, Italy):
- 5 L-Prolinamide,  
taurine,  
3-amino-1-propanesulfonic acid,  
3-aminopropylphosphonic acid,  
10 4-amino-1-benzylpiperidine.

4. Production of antibiotic GE 2270 for preparing antibiotic  
15 GE 2270 factors A, B<sub>1</sub>, B<sub>2</sub>, C<sub>1</sub>, C<sub>2</sub>, C<sub>2a</sub>, D<sub>1</sub>, D<sub>2</sub>, and E

A culture of Planobispora rosea ATCC 53773 is grown on an oatmeal agar slant for two weeks at 28-30°C and then used to inoculate 500 ml flasks containing  
20 100 ml of a seed medium of the following composition:

	Starch	20 g/l
	Polypeptone	5 g/l
25	Yeast extract	3 g/l
	Beef extract	2 g/l
	Soybean meal	2 g/l
	Calcium carbonate	1 g/l
30	Distilled water q.s.	100 ml
	(adjusted to pH 7.0 before sterilization)	

The flask is incubated on a rotary shaker (200 rpm) at 28-30°C for 92 h. The obtained culture is then used to inoculate a jar fermenter containing 4 liters of the same medium and the culture is incubated at 28-30°C for 48 hours with stirring (about 900 rpm) and

aeration (about one standard liter of air per volume per minute).

- 5           The obtained broth is transferred to a fermenter containing 50 l of the following production medium:

	Starch	20	g/l
10	Peptone	2.5	g/l
	Hydrolyzed casein	2.5	g/l
	Yeast extract	3	g/l
	Beef extract	2	g/l
	Soybean meal	2	g/l
15	Calcium carbonate	1	g/l
	Distilled water	q.s.	
	(adjusted to pH 7.0 before sterilization)		
	and incubated for about 72 hours at 28-30°C.		

- 20           Antibiotic production is monitored by paper disc agar assay using B. subtilis ATCC 6633 grown on minimum Davis medium. The inhibition zones are evaluated after incubation overnight at 35°C.

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4a)       Recovery of crude antibiotic GE 2270

- 30           The fermentation mass (50 l) obtained above is harvested and submitted to filtration in the presence of a filter aid (Clarcell).

Antibiotic GE 2270 is found mainly in the mycelium, even if a certain amount of it can be recovered also from the filtrates.



The filtrate is adjusted to about pH 7.0 and extracted with ethyl acetate (50 l). The organic phase is separated by centrifugation and concentrated to a small volume under reduced pressure. The obtained oily residue is then treated with petroleum ether to precipitate crude antibiotic GE 2270 that is collected by filtration and dried. 415 mg of crude antibiotic GE 2270 complex is obtained.

The mycelium is extracted twice with 20 l of methanol and the pooled extracts are concentrated under reduced pressure to give an aqueous residue which is extracted twice with ethyl acetate. Crude antibiotic GE 2270 (6.06 g) is precipitated by addition of petroleum ether from the concentrated organic phase.

4b) Isolation of antibiotic GE 2270 factor A

The crude obtained from the mycelium according to the procedure described above (3 g) is dissolved in tetrahydrofuran and concentrated under reduced pressure in the presence of silica gel (230-400 mesh). The obtained solid residue is collected and applied to a chromatography column containing 300 g of silica gel (230-400 mesh) prepared in methylene chloride ( $\text{CH}_2\text{Cl}_2$ ). The column is developed first with methylene chloride (2 l) and then sequentially with 1.5 l mixtures of methylene chloride and methanol in the following ratios: 98/2; 96/4, 94/6, 92/8, 90/10 and 88/12 (v/v).

Fractions are collected, analyzed by TLC, HPLC or microbiologically against B. subtilis and pooled according to their antibiotic content.

5

The pooled fractions containing antibiotic GE 2270 factor A are concentrated under reduced pressure to give an oily residue which is solubilized with tetrahydrofuran.

10

From this solution, antibiotic GE 2270 factor A (600 mg) is precipitated by adding petroleum ether.

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4bis) Isolation of mixtures of minor components of antibiotic GE 2270

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A representative mixture particularly enriched in the minor components C<sub>2a</sub>, D<sub>1</sub>, D<sub>2</sub> and E was established by HPLC comparison with analytical samples of each single component.

25

R<sub>t</sub> (min) refer to HPLC method M reported in the HPLC analysis section and are 20.55 for GE 2270 factor C<sub>2a</sub>, 17.43 for GE 2270 factor D<sub>1</sub>, 18.17 for GE 2270 factor D<sub>2</sub>, and 16.61 for GE 2270 factor E.

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Concentration of this fraction under reduced pressure produced an oily residue which was redissolved in tetrahydrofuran and precipitated with petroleum ether as whitish powder.

4c) Separation and isolation of antibiotic GE 2270 factors B<sub>1</sub>, B<sub>2</sub>, C<sub>1</sub>, C<sub>2</sub>, D<sub>1</sub>, D<sub>2</sub>, and E

5 Antibiotic GE 2270 factors D<sub>1</sub>, D<sub>2</sub> and E are separated and purified from the above obtained crude mixture by preparative HPLC using a 250x20 mm column packed with Nucleosil® C18 (silica gel functionalized with octadecylsilane groups) (5 µm) and eluted with  
10 mixtures of Phase A: CH<sub>3</sub>CN:tetrahydrofuran:40 mM HCOONH<sub>4</sub> (40:40:20); Phase B: CH<sub>3</sub>CN:tetrahydrofuran:40 mM HCOONH<sub>4</sub> (10:10:80). The antibiotic mixture (6 mg) was solubilized in 3 ml of Phase B and 1 ml of Phase A and  
15 was injected into the HPLC column which was eluted at a flow rate of 14 ml/min with a 26:74 mixture of Phase A and B. The eluted fractions were collected according to the UV adsorption profile at 254 nm. The fractions of subsequent chromatographic runs having homogeneous  
20 content were pooled and concentrated under reduced pressure to eliminate CH<sub>3</sub>CN. The residual solution showed antibacterial activity against Staphylococcus aureus Tour L165 by paper disc assay. These solutions were lyophilized at least three times to remove completely  
25 the HCOONH<sub>4</sub> buffer residue from the HPLC phases.

The yields were as follows: antibiotic GE 2270 factor E, 11 mg; antibiotic GE 2270 factor D<sub>1</sub>, 12 mg; antibiotic GE 2270 factor D<sub>2</sub>, 10 mg.

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4d) Isolation of a purified mixture containing antibiotic GE 2270 factor C<sub>2</sub> in mixture with other GE 2270 factors

The preparations of crude GE 2270 factors from 6 repeated fermentations were pooled and solubilized

into 12 l of  $\text{CH}_2\text{Cl}_2$ :methanol (93:7). The insoluble material was removed by filtration and the solution, containing the antibiotic complex, was applied to a 13 kg (230-400 mesh) silica gel column equilibrated in  $\text{CH}_2\text{Cl}_2$ :methanol (93:7). Antibiotic GE 2270 factor  $\text{C}_{2a}$  was eluted from the column by eluting with  $\text{CH}_2\text{Cl}_2$  : methanol (93:7). The fractions containing the antibiotic of the invention (HPLC analysis) were pooled, were concentrated under reduced pressure and were dried to yield 23.5 g of antibiotic GE 2270 factor  $\text{C}_{2a}$  in mixture with other minor factors.

A portion (5.5 g) of this preparation was again purified by flash chromatography on a column containing 400 g of silica gel (230-400 mesh) equilibrated in methylene chloride ( $\text{CH}_2\text{Cl}_2$ ). The column was developed first with methylene chloride (1 liter) and then sequentially with a series of mixtures of methylene chloride / methanol in the following ratios (v/v): 96/4 (3 liters); 94/6 (1 liter); 92/8 (2 liters); 90/10 (6 liters) and 88/12 (4 liters).

The fractions containing mainly GE 2270 factor  $\text{C}_{2a}$  (HPLC analysis) were pooled and were concentrated. The antibiotic preparation (646 mg) was precipitated upon addition of petroleum ether.

4e) Isolation of pure antibiotic GE 2270 factor  $\text{C}_{2a}$

The purified mixture containing mainly antibiotic GE 2270 factor  $\text{C}_{2a}$  was further purified by preparative HPLC from the above described preparation.

A portion of the above described preparation of the antibiotic (10 mg) was solubilized in 1 ml of Phase A ( $\text{CH}_3\text{CN}$  : tetrahydrofuran : 40 mM  $\text{HCOONH}_4$  - 40:40:20)

and 1 ml of Phase B ( $\text{CH}_3\text{CN}$  : tetrahydrofuran : 40 mM  $\text{HCOONH}_4$  -10:10:80) and was injected into a HPLC 250x20 mm Hibar column (E. Merck; Darmstadt F.R. Germany) packed with 7  $\mu\text{m}$  Nucleosil<sup>®</sup>Cl8 (silica gel functionalized with octadecylsilane groups) which was equilibrated with a mixture of 40% Phase A and 60% Phase B. The column was eluted at 15 ml/min flow rate with a 22 minutes linear gradient from 40% to 50% of Phase A. The UV detection was 254 nm. The fractions of 10 subsequent chromatographic runs containing the pure antibiotic of the invention were pooled and were concentrated under reduced pressure to eliminate  $\text{CH}_3\text{CN}$ . Antibiotic GE 2270 factor  $\text{C}_{21}$  precipitated from water. The precipitate was collected by centrifugation, was washed twice with distilled water and was dried under vacuum yielding 66 mg of the pure antibiotic.

20

#### 5. Preparation of antibiotic GE 2270 factor $\text{A}_2$

Antibiotic GE 2270 factor A (prepared as described above) (86 mg) is dissolved in 17 ml of 95% ethanol and 1.7 ml of acetic acid. After incubation at 60°C for 24 h, the resulting solution is diluted with 0.1M sodium phosphate buffer pH 7.5 (100 ml) and adjusted to pH 7.5 with 1M sodium hydroxide. Ethanol is removed by evaporation under reduced pressure and the aqueous residue is extracted twice with ethyl acetate (100 ml). The organic phase is concentrated under reduced pressure to obtain a solid residue which is solubilized with tetrahydrofuran and then precipitated by adding petroleum ether. Antibiotic GE 2270 factor  $\text{A}_2$  (62 mg) is obtained with minor amounts of antibiotic

GE 2270 factors A and A<sub>1</sub>. Pure antibiotic GE 2270 factor A<sub>2</sub> is obtained by preparative HPLC as follows:

5           10 Mg of the above crude product is solubilized  
in tetrahydrofuran, diluted to the solubility limit with  
water and then injected into a HPLC system with a column  
(250 x 20 mm) packed with Nucleosil<sup>®</sup> C18 (5 micrometer)  
reverse phase silica gel by Stacroma<sup>®</sup>, eluting with a  
10 linear gradient from 64% to 93% of phase B in phase A,  
in 20 min, at a flow rate of about 15 ml/min. In this  
system, phase A is a 90:10 (v/v) mixture of 18 mM sodium  
phosphate pH 7.2 and acetonitrile, while phase B is a  
15 40:60 (v/v) mixture of 18 mM sodium phosphate pH 7.2 and  
acetonitrile. Fractions of five consecutive runs are  
collected and UV monitored at 330 nm. Fractions which  
contain substantial amounts of antibiotic GE 2270 factor  
A<sub>2</sub>, which correspond to the major peaks of the UV  
20 elution profile, are pooled and concentrated under  
reduced pressure to an aqueous phase which is extracted  
twice with ethyl acetate. This organic layer is then  
washed with distilled water to remove the residual  
inorganic salts and concentrated to precipitate a solid  
25 residue that is then dissolved in tetrahydrofuran and  
re-precipitated with petroleum ether, to obtain pure  
antibiotic GE 2270 factor A<sub>2</sub> (45 mg).

30           In European Patent Application Publication  
No. 406745 are described other alternative method for  
preparing antibiotic GE 2270 factor A<sub>2</sub> as main reaction  
product of antibiotic GE 2270 factor A.

6. Preparation of antibiotic GE 2270 factor A<sub>3</sub>

Antibiotic GE 2270 factor A<sub>2</sub> is incubated for 1  
5 h at room temperature in 0.5M sodium carbonate. The  
reaction mixture is then diluted with cold water and  
brought to pH 6.5 with hydrochloric acid. The  
neutralized solution contains antibiotic GE 2270 factor  
A<sub>3</sub> as the main reaction product. This antibiotic is  
10 extracted from the aqueous phase with ethyl acetate and  
then is precipitated from the concentrated organic phase  
by adding petroleum ether.

Pure antibiotic GE 2270 factor A<sub>3</sub> is obtained by  
15 column chromatography as described below:

1.5 Grams of crude GE 2270 A<sub>3</sub> is dissolved in 60  
ml of a 1/1 (v/v) mixture of methanol and  
20 dichloromethane and adsorbed on silica gel (75-230 mesh)  
by evaporation of the solvents under reduced pressure.  
The solid residue is then put on the top of a silica gel  
(75-230 mesh) column (bed height 40 cm) equilibrated  
with dichloromethane. The column is then eluted with  
25 mixtures of methanol in dichloromethane in the order: 1)  
2% methanol (450 ml); 2) 5% methanol (500 ml); 3) 10%  
methanol (600 ml); 4) 15% methanol (500 ml); 5) 20%  
methanol (500 ml); 6) 30% methanol (250 ml).

Fractions are collected and monitored by TLC and  
30 a microbiological assay on B. subtilis ATCC 6633.  
Antibiotic GE 2270 factor A<sub>3</sub> is normally present in the  
eluates which contain about 15-20% methanol.

The fractions containing the desired product are pooled and concentrated under reduced pressure. Upon addition of petroleum ether to the residue, antibiotic  
5 GE 2270 factor A<sub>3</sub> precipitates (854 mg of pure product).

10 7. Preparation of the proper starting material from antibiotic factors D<sub>1</sub>, D<sub>2</sub>, E and C<sub>2a</sub>

By substantially following the same procedure described at points 5 and 6 above but starting from the  
15 single factors D<sub>1</sub>, D<sub>2</sub>, E and C<sub>2a</sub> of antibiotic GE 2270 instead of factor A, the proper starting materials of formula III wherein W is COOH or an activated ester, R is hydrogen or CH<sub>2</sub>OH, R<sub>1</sub> is CH<sub>3</sub> or hydrogen and R<sub>4</sub> is  
20 hydroxymethyl or methyl, are obtained.

7a) Preparation of proper starting material from a mixture of minor components (C<sub>2a</sub>, D<sub>1</sub>, D<sub>2</sub> and E) of antibiotic GE 2270  
25

By substantially following the same procedure described at point 5 and 6 above but starting from a mixture of minor components (C<sub>2a</sub>, D<sub>1</sub>, D<sub>2</sub> and E) of  
30 antibiotic GE 2270 instead of the single factor A, the proper starting material of formula III wherein W is COOH or an activated ester and R, R<sub>1</sub> and R<sub>4</sub> are respectively methoxymethyl, methyl and hydroxymethyl for C<sub>2a</sub>, hydrogen, hydrogen and methyl for D<sub>1</sub>, hydroxymethyl, methyl and methyl for D<sub>2</sub> and hydroxymethyl, hydrogen and methyl for E are obtained.



$R_t$  (min) refer to HPLC method M reported in the HPLC analysis section.

5 When W is an activated ester,  $R_t$  (min) are respectively 22.51 for GE 2270 factor  $C_{2a}$ , 19.80 for GE 2270 factor  $D_1$ , 20.41 for GE 2270 factor  $D_2$  and 18.92 for GE 2270 factor E.

10 When W is COOH,  $R_t$  (min) are respectively 12.99 for GE 2270 factor  $C_{2a}$ , 10.38 for GE 2270 factor  $D_1$ , 11.08 for GE 2270 factor  $D_2$  and 9.03 for GE 2270 factor E.

15 8. Preparation of glycyl-N $\epsilon$ -Cbz-L-lysine trifluoroacetate

4.8 ml of DPPA (22 mmols) was added at 0°C to a well stirred solution of 3.5 g of BOC-glycine (Fluka) (20 mmols) and 7.28 g of N $\epsilon$ -Cbz-L-lysine methyl ester hydrochloride (Fluka) (22 mmols) in 50 ml of dry DMF. To this solution, a solution of 5.8 ml of TEA (42 mmols) in 50 ml of dry DMF was added at 0°C over a 10 - 15 min period. Stirring was continued for 2 more hours at 0°C and then overnight at room temperature. 25 The reaction mixture was diluted with 250 ml of toluene and 500 ml of ethyl acetate and washed with 1N aq. HCl (x3), water, a saturated solution of NaHCO<sub>3</sub> and brine. Drying over Na<sub>2</sub>SO<sub>4</sub> and evaporation of the solvent yielded 9.7 g of a thick oil which resisted any attempt 30 of crystallization. NMR of this oil was in perfect agreement with the structure of BOC-glycyl-N $\epsilon$ -Cbz-L-lysine methyl ester.

The oil was dissolved in 200 ml of acetone/dioxane 1:1 and 22 ml of 1N aq. NaOH were added over a 30 min period at 0°C under stirring. The

reaction was then stirred for 45 min at room temperature, diluted with 300 ml of cold water, acidified with 25 ml of 1N aq HCl and extracted with chloroform (x3) and ethyl acetate (x3). Drying over Na<sub>2</sub>SO<sub>4</sub> and evaporation of the solvent yielded 9.4 g of a gum which resisted any attempt of crystallization. NMR of this gum was in perfect agreement with the structure of BOC-glycyl-N $\epsilon$ -Cbz-L-lysine.

10

The gummy compound was treated with 20 ml of cold trifluoroacetic acid (TFA). The reaction mixture was swirled at room temperature until all the compound went in solution. The solution was reduced to a small volume under vacuum in the cold and then ethyl ether was added to induce precipitation of the title compound. 9.6 g of glycyl-N $\epsilon$ -Cbz-L-lysine trifluoroacetate were obtained as a white powder. NMR was in perfect agreement with the structure.

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#### 9. Preparation of L-tyrosyl-L-prolinamide

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0.48 ml of DPPA (2 mmols) were added at 0°C to a well stirred solution of 538 mg of BOC-L-tyrosine (Fluka) (2 mmols), 228.3 mg of L-prolinamide (Aldrich) (2 mmols) and 168 mg of NaHCO<sub>3</sub> in 5 ml of dry DMF. The reaction was stirred for 24 h at room temperature and then diluted with 50 ml of water and extracted with chloroform (x3). The organic phase was washed water, dried over Na<sub>2</sub>SO<sub>4</sub> and the solvent evaporated to yield an oil which was purified by flash chromatography on silica gel 60 (230 - 400 mesh ASTM - Merck) eluting with hexane/acetone 2:3. 420 mg of BOC-L-tyrosyl-L-

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prolinamide were in this way obtained as a white solid. NMR was in agreement with the structure.

5           The solid obtained was dissolved in 6 ml of ethyl acetate and stirred for 48 h at room temperature in the presence of 4 ml of 3N aq. HCl. The reaction mixture was then evaporated to dryness in vacuo and the residue redissolved in ethanol was precipitated with  
10 ethyl ether. 302 mg of L-tyrosyl-L-prolinamide were obtained as a white powder. NMR was in perfect agreement with the structure.

15

10.       Preparation of methyl 8-aminocaprylate and methyl 11-aminoundecanoate p-toluenesulfonates

20           A solution of 40 mmols of the selected amino acid (Fluka) and 15.2 g of p-toluenesulfonic acid monohydrate (Fluka) (80 mmols) in 200 ml of methanol was refluxed overnight. The solvent was then evaporated in vacuo and the residue redissolved in ethyl ether.  
25 After sometime the title compounds crystallized out quantitatively. The NMR of both compounds was in agreement with their structure.

30

11.       Preparation of 5-aminopentylphosphonic acid

3.48 g of 5-amino-1-pentanol (Fluka) (33.7 mmols) and 5.0 g of phthalic anhydride (Fluka) (33.7 mmols) were melted together at 180°C. This temperature was maintained for 90 min until no more

water developed. The reaction was allowed to cool to room temperature and the oily mixture was chromatographed on silica gel 60 (230 - 400 mesh ASTM - Merck) eluting with 2% methanol in chloroform. 5.9 g of a pure oil were obtained. NMR was in agreement with the structure.

To the 5.9 g of the oily intermediate (25 mmols), 1.6 ml of  $\text{PBr}_3$  (17 mmols) were added portionwise so to control the exothermic reaction. The reaction mixture was heated at  $100^\circ\text{C}$  for 1.5 h and then poured into crushed ice. The solid material that separated was filtered and allowed to dry in air overnight. 6.6 g of the pure bromo intermediate were obtained. The mass was in agreement with the expected molecular weight.

500 mg of the pure bromo intermediate (1.69 mmols) and 140 mg of triethyl phosphite (Fluka) (0.84 mmols) were heated together at  $150^\circ\text{C}$  for about 1 h. Other three portions of 140 mg of triethyl phosphite were then added at 30 min interval at the same temperature. When all the starting material had disappeared, the excess of triethyl phosphite was distilled off and the crude material purified by flash chromatography on silica gel 60 (230 - 400 mesh ASTM - Merck) eluting with 2% methanol in dichloromethane. 468 mg of the expected diethyl phosphonate were obtained as a thick oil. NMR confirmed the structure.

468 mg of the diethyl phosphonate intermediate were treated overnight with 3 ml of a 0.2 M solution of hydrazine in methanol at room temperature. The precipitated phthalhydrazide was filtered off and the

remaining solution was evaporated to dryness in vacuo. The residue was taken up in 1 N aq. HCl and the solution was washed with ethyl acetate, basified with NaOH and extracted several times with n-butanol. The butanolic phase was dried over Na<sub>2</sub>SO<sub>4</sub> and evaporated to dryness to yield 175 mg of a thick oil whose NMR was in agreement with the product expected.

175 mg of diethyl 5-aminopentylphosphonate were refluxed for 20 h in 0.6 ml of conc. HCl. The acid solution was then evaporated to dryness by azeotropic distillation in vacuo in the presence of n-butanol. The NMR of the glassy oil obtained confirmed it to be the 5-aminopentylphosphonic acid.

#### Preparation of 5-(5-aminopentyl)tetrazole

To a solution of 10 ml of 6-aminocapronitrile (Fluka) (80 mmols) and 13.3 ml of TEA (96 mmols) in 80 ml of tetrahydrofuran, 12.48 ml of benzyl chloroformate (Fluka) (88 mmols) were added dropwise at 0°C under stirring. Stirring was continued for 2 h at room temperature and the solvent was evaporated in vacuo. The residue was dissolved in ethyl acetate, washed with 1 N aq. HCl, water and then dried over Na<sub>2</sub>SO<sub>4</sub> and the solvent evaporated to yield 19.6 g of a syrup whose NMR was in agreement with the structure.

1 g of the protected 6-aminocapronitrile (4.06 mmols) in 40 ml of 1-methyl-2-pyrrolidone was heated at 150°C under argon in the presence of 793 mg of sodium azide (12.2 mmols) and 834 mg triethylamine

hydrochloride (6.1 mmols). After 4 h the reaction mixture was diluted with 120 ml of water and then carefully acidified to pH 1 with 10% aq. HCl (attention: azotidric acid forms!). The solution was extracted with ethyl acetate, the organic phase re-extracted with 10% aq. NaOH (x2) and the basic solution washed with ethyl ether, acidified with conc. HCl and extracted with ethyl acetate (x3). Drying and evaporation of the organic phase yielded a syrup that crystallized from methanol/water. 260 mg of a fine powder were obtained. NMR and mass confirmed the structure.

250 mg of the N-protected amino tetrazole (0.86 mmols) were treated at room temperature with 5ml of thioanisole (43.25 mmols) and 17.5 ml of trifluoroacetic acid for 3 h. Trifluoroacetic acid was concentrated in vacuo in the cold and ethyl ether was added to precipitate the title compound as its trifluoroacetate salt. NMR and mass confirmed the structure.

25

13. Preparation of N-[3,4-di-(O-tetrahydropyranyl)benzoyl]-thiazolidin-2-thione

A solution of 4.62 g of 3,4-dihydroxybenzoic acid (Fluka) (30 mmols) in 40 ml of methanol was refluxed for 24 h in the presence of 0.325 ml of conc. H<sub>2</sub>SO<sub>4</sub>. After cooling the solution to room temperature some solid NaHCO<sub>3</sub> was added and the solvent evaporated in vacuo. The residue was taken up in ethyl acetate, washed with water, dried over Na<sub>2</sub>SO<sub>4</sub> and the solvent

evaporated to yield a syrup which was crystallized from ethyl acetate/hexane. 3.53 g of white crystals were obtained.

5

9.1 ml of dihydropyran (Fluka) (0.1 mol) and 250 mg of pyridinium p-toluenesulfonate (1 mmol) were added at room temperature to a stirred solution of 1.68 g of methyl 3,4-dihydroxybenzoate (10 mmols) in 4 ml of ethyl acetate and 25 ml of dichloromethane. After 4 d the reaction mixture was washed with a saturated solution of  $\text{NaHCO}_3$ , dried over  $\text{Na}_2\text{SO}_4$  and evaporated to dryness to obtain 3.36 g of an oil which was used for the next step without further purification.

15

The crude from the previous reaction was dissolved in 40 ml of acetone and to the stirred solution 20 ml of water, 2.76 g of  $\text{K}_2\text{CO}_3$  (20 mmols) and 10 ml of 1N aq.  $\text{NaOH}$  (10 mmols) were added and stirring was continued for 7 d at room temperature. Acetone was evaporated in vacuo and the residual water phase was washed with ethyl acetate. The aqueous phase was transferred to an E. flask containing an equal volume of chloroform, cooled to  $0^\circ\text{C}$  and carefully acidified under vigorous stirring with 50 ml of 1 N aq.  $\text{HCl}$ . The water phase was then extracted 3 more times with chloroform and the combined organic layers were washed with 0.2% ammonium formate, dried over  $\text{Na}_2\text{SO}_4$  and evaporated to dryness to yield a syrup which crystallized after hexane addition. 2.34 g of a white solid were obtained. The NMR was in agreement with the structure.

25  
30

333 mg of 2-thiazoline-2-thiol (Fluka) (2.8 mmols), 577 mg of N,N'-dicyclohexylcarbodiimide

(Fluka) (2.8 mmols) and 35 mg of 4-dimethylamino-pyridine were added in the order at 0°C to a stirred solution of 644 mg of the benzoic acid intermediate (2 mmols) in 14 ml ethyl acetate/dichloromethane 5:2. 5  
Stirring was continued overnight at room temperature, the precipitated dicyclohexylurea was filtered off and the yellow solution was evaporated in vacuo to yield a yellow oil which was purified by flash chromatography 10  
on silica gel 60 (230 - 400 mesh ASTM - Merck) eluting from 25% acetone in hexane. 700 mg of yellow crystals were obtained from acetone/hexane. NMR and IR confirmed the compound to be the title compound.

15

14. Preparation of N1-N8-di-tert-butoxycarbonylspermidine

20 A solution of 19.72 g of BOC-ON (Aldrich) (80 mmols) in 60 ml of degassed tetrahydrofuran (THF) was added dropwise over a 1 h period under argon to a stirred solution of 5.8 g of spermidine (Aldrich) (40 mmols) in 40 ml of degassed THF cooled at 0°C. The 25  
reaction was then stirred at room temperature overnight and then evaporated to dryness. The residue was taken up in ethyl ether, washed with 1 N aq. NaOH (x4) and water (x4), dried over Na<sub>2</sub>SO<sub>4</sub> and the solvent concentrated to a small volume in vacuo. Upon addition 30  
of ethyl ether 11 g of a white powder precipitated. NMR confirmed it to be the title compound.



15. Preparation of N-tert-butoxycarbonylpropilendiamine

4.2 g of BOC-ON (Aldrich) (17.2 mmols) were  
5 added at room temperature to a stirred solution of 2 g  
of 3-aminopropionitrile fumarate (Aldrich) (15.6 mmols)  
dissolved in a mixture of 10 ml of dioxane, 10 ml of  
water and 3.3 ml of triethylamine. After 3 h the  
reaction mixture was diluted with more water and  
10 extracted with dichloromethane (x3). The combined  
organic layers were washed with 1 N aq. NaOH (x3) and  
water (x3), dried over  $\text{Na}_2\text{SO}_4$  and evaporated to  
dryness. The residual oil was taken up in ethyl ether  
and precipitated with hexane to yield 2.2 g of a white  
15 powder.

1 g of N-BOC-protected intermediate (5.9 mmols)  
in 7 ml of 1 N ethanolic NaOH was hydrogenated at 40  
20 psi in the presence of 130 mg of Raney nickel (50%  
slurry in water,  $\text{pH} > 9$ ) (Aldrich) for 40 h. Raney nickel  
was filtered off and the solvent was evaporated to  
dryness. The residue was taken up in ethyl acetate and  
washed with 1 N aq. NaOH, dried over  $\text{Na}_2\text{SO}_4$  and the  
25 solvent removed in vacuo yielding 950 mg of a colorless  
oil which solidified on standing. NMR confirmed it to  
be the title compound.

30 16. Preparation of 3-(2-aminoethylthio)propanoic acid methyl  
ester trifluoroacetate

To a solution of 0.5 g of cysteamine (Fluka)  
(6.48 mmols) in 5 ml of  $\text{CH}_2\text{Cl}_2$ , 1.4 g of di-tert-butyl  
dicarbonate (Aldrich) (6.48 mmols) in 5 ml of  $\text{CH}_2\text{Cl}_2$   
were added at room temperature under stirring. After 30

min the organic solvent was evaporated and the crude material dissolved in 5 ml of absolute ethanol.

To the ethanolic solution, 2.7 ml of TEA (19.1 mmols) and 1.07 ml of methyl 3-bromopropionate (Fluka) (9.57 mmols) were added in the order. The reaction was completed in about 30 min. Ethanol was removed in vacuo and replaced by 15 ml of chloroform. The organic phase was then washed with water, anidrified on Na<sub>2</sub>SO<sub>4</sub> and the solvent evaporated to yield an oil which was finally treated with 1 ml of trifluoroacetic acid at 0°C for 5 min. Evaporation to dryness gave 270 mg of a pale yellow oil. NMR and IR confirmed it to be the title compound.

#### 17. Preparation of 6-amino-2(E)-hexenoic acid

To a stirred solution of 2 ml of 4-amino-butyraldehyde diethyl acetal (Fluka) (11.6 mmols) and 3.6 ml of TEA (25.6 mmols) in 5 ml of CH<sub>2</sub>Cl<sub>2</sub>, a solution of 1.5 ml of benzoyl chloride (Fluka) (12.9 mmols) in 5 ml CH<sub>2</sub>Cl<sub>2</sub> was added in 30 min. at room temperature. After 1 hour the reaction was diluted with 10 more ml of CH<sub>2</sub>Cl<sub>2</sub>, washed with water and the organic phase dried over Na<sub>2</sub>SO<sub>4</sub> and the volume adjusted to 20 ml. The new solution was allowed to react for three days under argon in the presence of 1.6 ml of TEA (11.5 mmols), 10.2 g of di-tert-butyl dicarbonate (Aldrich) (46.8 mmols) and 1.4 g of 4-dimethylamino-pyridine (Fluka) (11.5 mmols) at room temperature. Removal of the solvent gave a brown oil that was purified by flash chromatography on silica gel 60 (230 - 400 mesh ASTM - Merck) eluting with 20% ethyl

acetate in n-hexane yielding 1.6 g of the N,N deprotected 4-aminobutyraldehyde diethyl acetal as a colorless oil. NMR confirmed the structure.

5

The obtained oil was then dissolved in 5 ml of THF and treated with 5 ml of 1N HCl at room temperature for three hours. THF was removed in vacuo and the remaining solution was washed with chloroform (2 ml x3).

10

The organic phase was then washed with a solution of  $\text{Na}_2\text{CO}_3$ , water, dried over  $\text{Na}_2\text{SO}_4$  and evaporated to dryness yielding an oil that was used in the next step without further purification.

15

To a suspension of 160 mg of 60% NaH (4mmols) in 5 ml of dry THF at 0°C under argon, 0.837 ml of triethylphosphonoacetate (Fluka) (4.3 mmols) were added. After 30 min. a dry THF (2 ml) solution of the

20

previously obtained aldehyde (1.17 g) (4.02 mmols) was added and the temperature was allowed to rise to room temperature. The reaction was stirred overnight and then 50 more mg of 60% NaH were added at 0 C. After two more hours at room temperature the reaction mixture was

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treated with diluted HCl (10 ml) and extracted with ethyl acetate (5ml x3). The combined organic phase was washed with water, dried over  $\text{Na}_2\text{SO}_4$  and evaporated to dryness. The crude material was purified by flash

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chromatography on silica gel 60 (230 - 400 mesh ASTM - Merck) eluting with 15% ethyl acetate in n-hexane yielding 765 mg of a syrup. NMR confirmed it to be the expected product with the double bond in E configuration ( $J = 16\text{Hz}$ ).

6.15 ml of 1N LiOH (6.15 mmols) were added to a solution of 739 mg of the unsaturated ester previously

obtained (2.05 mmols) in 10 ml of THF under stirring at room temperature. When the starting material had disappeared the reaction mixture was concentrated in vacuo at 30°C (bath temperature). The aqueous solution was acidified at pH 2 with 1N HCl and then extracted with ethyl acetate. The combined organic phase was dried over Na<sub>2</sub>SO<sub>4</sub>, filtered and the solvent evaporated yielding an oil that solidified upon standing under vacuum. NMR and MS confirmed it to be 6-N-BOC-amino-2(E)-hexenoic acid.

Removal of the N-BOC protection to obtain the title compound was carried out in neat trifluoroacetic acid at 0°C just before the coupling with the appropriate GE 2270 starting material.

18. Preparation of 3-(2-aminoethoxy)propanoic acid trifluoroacetate

To a stirred solution of 1 g of N-BOC-ethanolamine (6.22 mmols) [prepared according to classical methodologies from ethanolamine (Fluka)] in 10 ml of dry THF at -78 C, 3.88 ml of 1.6 M solution of butyllithium (Fluka) (6.22 mmols) were added under argon. After 30 min. 1.3 g of t-butyl 3-bromo propanoate [prepared according to classical methodologies from 3-bromo propanoic acid (Fluka)] (6.22 mmols) were added, the temperature allowed to rise to room temperature and the resulting mixture stirred for 20 hours at that temperature. After dilution with water the reaction mixture was extracted with n-hexane (5ml x2). Removal of the solvent gave a crude material that was purified by

flash chromatography on silica gel 60 (230 - 400 mesh  
ASTM - Merck) eluting with 20% ethyl acetate in n-hexane  
yielding 1.43 g of an oil. NMR confirmed it to be the  
coupled compound.

5

The total deprotection of the coupled compound  
was carried out immediately before addition to the  
appropriate GE 2270 starting material by stirring it in  
trifluoroacetic acid for about 5 min at room  
temperature. Removal of trifluoroacetic acid in vacuo  
yielded the title compound.

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CLAIMS

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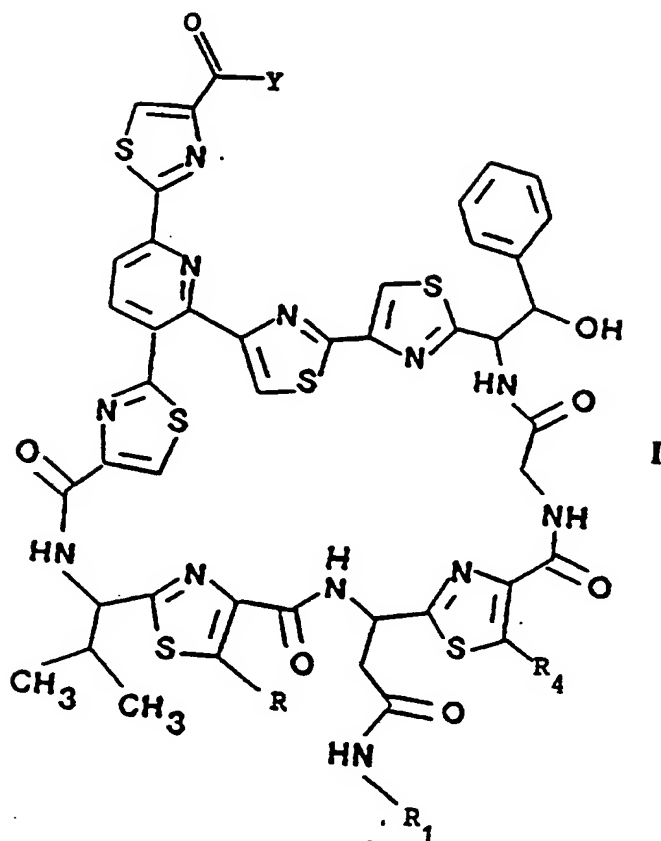
1. An amide derivative of antibiotic GE 2270  
10 having the following formula I

15

20

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30



wherein

R represents:  
hydrogen,  
hydroxymethyl, or  
5 methoxymethyl;

R<sub>1</sub> represents:  
hydrogen, or  
10 methyl;

Y represents:  
15 a group of formula



wherein:

R<sub>2</sub> represents:  
25 hydrogen,  
(C<sub>1</sub>-C<sub>4</sub>)alkyl,  
amino(C<sub>2</sub>-C<sub>4</sub>)alkyl,  
(C<sub>1</sub>-C<sub>4</sub>)alkylamino-(C<sub>1</sub>-C<sub>4</sub>)alkyl, or  
30 di-(C<sub>1</sub>-C<sub>4</sub>)alkylamino-(C<sub>1</sub>-C<sub>4</sub>)alkyl;

R<sub>3</sub> represents:  
hydrogen,  
a linear or branched (C<sub>1</sub>-C<sub>14</sub>)alkyl group bearing  
from 1 to 3 substituents selected from: carboxy,

5 sulfo, phosphono, amino which may be optionally  
protected with a lower alkoxy carbonyl or a  
benzyloxy carbonyl group, (C<sub>1</sub>-C<sub>4</sub>)alkylamino  
wherein the alkyl moiety may be optionally  
substituted with a carboxy group,  
10 di-(C<sub>1</sub>-C<sub>4</sub>)alkylamino, hydroxy, halo,  
(C<sub>1</sub>-C<sub>4</sub>)alkoxy wherein the alkyl moiety may be  
optionally substituted with a carboxy group,  
(C<sub>1</sub>-C<sub>4</sub>)alkoxy carbonyl, mercapto,  
15 (C<sub>1</sub>-C<sub>4</sub>)alkylthio wherein the alkyl moiety may be  
optionally substituted with a carboxy group,  
phenyl which may be optionally substituted with  
1 to 3 substituents selected from carboxy,  
sulfo, hydroxy, halo and mercapto, carbamyl,  
20 (C<sub>1</sub>-C<sub>6</sub>)alkylcarbamyl wherein the alkyl moiety  
may be optionally substituted with 1 or 2  
substituents selected from carboxy, amino,  
(C<sub>1</sub>-C<sub>4</sub>)alkylamino and di-(C<sub>1</sub>-C<sub>4</sub>)alkylamino,  
di-(C<sub>1</sub>-C<sub>4</sub>)alkylcarbamyl wherein the alkyl  
moieties together with the adjacent nitrogen  
atom may also represent a saturated 5-7 membered  
heterocyclic ring which may optionally be  
substituted with a carboxy or a carbamyl group  
25 on one of the ring carbons and may optionally  
contain a further heterogroup selected from O, S  
and N, benzoylamino wherein the phenyl group may  
be substituted from 1 to 3 hydroxy group, a  
nitrogen containing 5-6 membered heterocyclic  
30 ring which may be unsaturated, partially  
saturated or wholly saturated and may contain 1  
to 3 further heteroatoms selected from N, S and  
O wherein one of the carbons of the ring may  
optionally bear a group carboxy, sulfo,  
carboxy(C<sub>1</sub>-C<sub>4</sub>)alkyl and sulfo(C<sub>1</sub>-C<sub>4</sub>)alkyl and



the ring nitrogen atom may optionally be substituted by (C<sub>1</sub>-C<sub>4</sub>)alkyl, carboxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, sulfo(C<sub>1</sub>-C<sub>4</sub>)alkyl, and benzyl;

5 (C<sub>3</sub>-C<sub>6</sub>)alkenyl, optionally substituted by carboxy or sulfo;  
1-deoxy-1-glucityl;  
2-deoxy-2-glucosyl;  
10 a fully saturated 5 to 7 membered nitrogen containing heterocyclic ring wherein the nitrogen atom may be optionally substituted by (C<sub>1</sub>-C<sub>4</sub>)alkyl or benzyl and one or two carbons of the ring skeleton may bear a substituent  
15 selected from (C<sub>1</sub>-C<sub>4</sub>)alkyl, carboxy and sulfo;

or R<sub>2</sub> and R<sub>3</sub>

20 taken together with the adjacent nitrogen atom represent a fully saturated 5-7 membered heterocyclic ring which may optionally contain a further heteroatom selected from O, S and N, and may optionally bear one or two substituents on the ring carbons selected from (C<sub>1</sub>-C<sub>4</sub>)alkyl,  
25 benzyl, carboxy, sulfo, carboxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, and sulfo(C<sub>1</sub>-C<sub>4</sub>)alkyl;

R<sub>4</sub> represents:  
30 hydrogen,  
methyl, or  
hydroxymethyl;

with the proviso that when  $R_4$  is hydrogen or hydroxymethyl, then simultaneously R is methoxymethyl and  $R_1$  is methyl;

5 and the pharmaceutically addition salts thereof.

2. A compound according to claim 1 wherein R represents methoxymethyl and the other substituents are  
10 defined as in claim 1.

3. A compound as claimed in claim 1 wherein R represents methoxymethyl,  $R_1$  and  $R_4$  represent methyl and  
15 Y represents a group of formula



25 wherein  $R_2$  is hydrogen and  $R_3$  is defined as in claim 1.

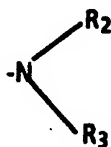
4. A compound as claimed in claim 1 wherein R is methoxymethyl,  $R_1$  and  $R_4$  represent a methyl group and Y  
30 is an amino moiety which derive from a natural amino acid such as for example glycine, ornithine, serine, aspartic acid, tyrosine, leucine, phenylalanine, methionine, proline, threonine, lysine, or a synthetic dipeptide such as glycyllysine, serylproline,

glycylprolinamide, tyrosylprolinamide,  
threonylprolinamide, leucylprolinamide.

5            5. A compound as claimed in claim 1 wherein R is  
methoxymethyl,  $R_1$  and  $R_4$  are methyl, Y is a group  $NR_2R_3$   
wherein  $R_2$  is hydrogen and  $R_3$  is a linear alkyl chain  
preferably of 3 to 12 carbons, more preferably of 3 to 7  
10 carbons substituted with a group selected from COOH,  
 $SO_3H$  and  $PO_3H_2$ .

6. A compound as claimed in claim 1 wherein R is  
15 methoxymethyl,  $R_1$  and  $R_4$  are methyl, Y is a group  $NR_2R_3$   
wherein  $R_2$  is hydrogen and  $R_3$  is  $CH_2CH_2CH_2CH_2COOH$ .

7. A compound as claimed in claim 1 wherein R  
20 represents hydrogen, hydroxymethyl or methoxymethyl,  $R_1$   
represents hydrogen or methyl,  $R_4$  represents hydrogen,  
methyl or hydroxymethyl and Y represents a group of  
formula



wherein  $R_2$  is hydrogen and  $R_3$  is defined as in  
claim 1.

8. A compound as claimed in claim 7 wherein Y is an amino moiety which derive from a natural amino acid such as for example glycine, ornithine, serine, aspartic acid, tyrosine, leucine, phenylalanine, methionine, proline, threonine, lysine, or a synthetic dipeptide such as glycylllysine, serylproline, glycyprolinamide, tyrosylprolinamide, threonylprolinamide, leucylprolinamide.

10

9. A compound as claimed in claim 1 wherein R is hydrogen, hydroxymethyl or methoxymethyl,  $R_1$  is hydrogen or methyl,  $R_4$  is hydrogen, methyl or hydroxymethyl and Y is a group  $NR_2R_3$  wherein  $R_2$  is hydrogen and  $R_3$  is a linear alkyl chain preferably of 3 to 12 carbons, more preferably of 3 to 7 carbons substituted with a group selected from COOH,  $SO_3H$  and  $PO_3H_2$ .

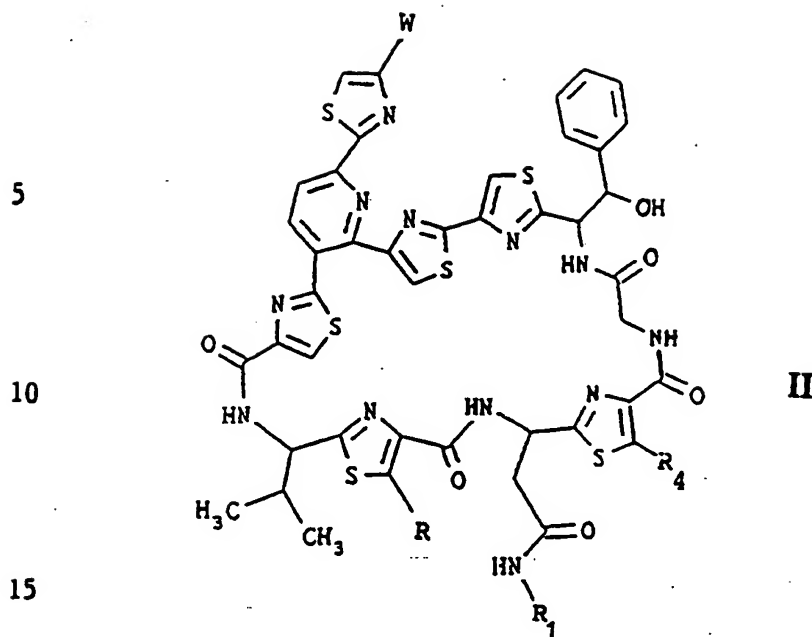
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10. A compound as claimed in claim 1 wherein R is hydrogen, hydroxymethyl or methoxymethyl,  $R_1$  is hydrogen or methyl,  $R_4$  is hydrogen, methyl or hydroxymethyl and Y is a group  $NR_2R_3$  wherein  $R_2$  is hydrogen and  $R_3$  is  $CH_2CH_2CH_2CH_2CH_2-COOH$ .

25

11. A process for preparing a compound of claim 1 which comprises reacting an antibiotic GE 2270 compound having formula II:

30



25 wherein

W represents a carboxy or an activated ester function;

R represents hydrogen, hydroxymethyl or methoxymethyl;

30 R<sub>1</sub> represents hydrogen or methyl;

R<sub>4</sub> represents hydrogen, methyl or hydroxymethyl;

with the proviso that when R<sub>4</sub> is hydrogen or hydroxymethyl, then simultaneously R is methoxymethyl and R<sub>1</sub> is methyl, with a selected amine of formula HNR<sub>2</sub>R<sub>3</sub> wherein R<sub>2</sub> and R<sub>3</sub> have the same meanings as in

claim 1, in an inert organic solvent and, when W is carboxy, in the presence of a condensing agent.

5           12. A process according to claim 11 wherein the condensing agents are selected from (C<sub>1</sub>-C<sub>4</sub>)alkyl, phenyl or heterocyclic phosphorazidates such as, diphenyl-phosphorazidate (DPPA), diethyl-phosphorazidate, di(4-nitrophenyl)phosphorazidate, dimorpholyl-  
10 phosphorazidate and diphenylphosphorochloridate.

          13. A process according to claims 11 and 12  
15 wherein the amine reactant HNR<sub>2</sub>R<sub>3</sub> is used in a 1 to 2 fold molar excess with respect on the antibiotic starting material and the reaction temperature is comprised between 0 and 20°C.

20           14. A compound of any of claims 1 to 10 for use as a medicine.


25           15. A pharmaceutical composition containing a compound of any of claims 1 to 10 as the active ingredient in admixture with a pharmaceutically acceptable carrier.

30           16. Use of a compound according to any of claims 1 to 10 for preparing a medicament for use as an antibiotic.

## INTERNATIONAL SEARCH REPORT

PCT/EP 92/00002

International Application

<b>I. CLASSIFICATION OF SUBJECT MATTER</b> (If several classification symbols apply, indicate all) <sup>6</sup>		
According to International Patent Classification (IPC) or to both National Classification and IPC		
Int.Cl. 5 C07K7/56; C07K5/06; A61K35/66		
<b>II. FIELDS SEARCHED</b>		
Minimum Documentation Searched <sup>7</sup>		
Classification System	Classification Symbols	
Int.Cl. 5	C07K ; A61K	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched <sup>8</sup>		
<b>III. DOCUMENTS CONSIDERED TO BE RELEVANT<sup>9</sup></b>		
Category <sup>10</sup>	Citation of Document, <sup>11</sup> with indication, where appropriate, of the relevant passages <sup>12</sup>	Relevant to Claims No. <sup>13</sup>
	No relevant documents disclosed  ---	
<p><sup>10</sup> Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"A" document member of the same patent family</p>		
<b>IV. CERTIFICATION</b>		
Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report	
18 MARCH 1992	27. 03. 92	
International Searching Authority	Signature of Authorized Officer	
EUROPEAN PATENT OFFICE	KORSNER S.E. 	

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